

PCTWORLD INTELLECTUAL PROPERTY ORGANIZATION
International Bureau

INTERNATIONAL APPLICATION PUBLISHED UNDER THE PATENT COOPERATION TREATY (PCT)

(51) International Patent Classification ⁶ : A61K 7/13, 7/00	A1	(11) International Publication Number: WO 99/13847 (43) International Publication Date: 25 March 1999 (25.03.99)
(21) International Application Number: PCT/US97/16578 (22) International Filing Date: 17 September 1997 (17.09.97) (71) Applicant (for all designated States except US): THE PROCTER & GAMBLE COMPANY [US/US]; One Procter & Gamble Plaza, Cincinnati, OH 45202 (US). (72) Inventors; and (75) Inventors/Applicants (for US only): MITSUMATSU, Arata [JP/JP]; 6-16-306, Midorigaoka-cho, Ashiya-shi, Hyogo 659 (JP). SALVADOR, Dorthy, Yong, Juanico [PH/JP]; #201, 9-1, Kusunoki-cho, Ashiya-shi, Hyogo 659 (JP). (74) Agents: REED, T., David et al.; The Procter & Gamble Company, 5299 Spring Grove Avenue, Cincinnati, OH 45217 (US).		(81) Designated States: AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ, DE, DK, EE, ES, FI, GB, GE, GH, HU, ID, IL, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK, MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZW, ARIPO patent (GH, KE, LS, MW, SD, SZ, UG, ZW), Eurasian patent (AM, AZ, BY, KG, KZ, MD, RU, TJ, TM), European patent (AT, BE, CH, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE), OAPI patent (BF, BJ, CF, CG, CI, CM, GA, GN, ML, MR, NE, SN, TD, TG). Published <i>With international search report.</i>
(54) Title: HAIR CARE COMPOSITIONS COMPRISING OPTICAL BRIGHTENERS AND NON-VOLATILE SOLVENTS (57) Abstract Disclosed are hair care compositions comprising: (a) an effective amount of an optical brightener; and (b) a non-volatile solvent for solubilizing at least 0.01 wt.% of the optical brightener, wherein the optical brightener is deposited on the hair without crystallizing.		

FOR THE PURPOSES OF INFORMATION ONLY

Codes used to identify States party to the PCT on the front pages of pamphlets publishing international applications under the PCT.

AL	Albania	ES	Spain	LS	Lesotho	SI	Slovenia
AM	Armenia	FI	Finland	LT	Lithuania	SK	Slovakia
AT	Austria	FR	France	LU	Luxembourg	SN	Senegal
AU	Australia	GA	Gabon	LV	Latvia	SZ	Swaziland
AZ	Azerbaijan	GB	United Kingdom	MC	Monaco	TD	Chad
BA	Bosnia and Herzegovina	GE	Georgia	MD	Republic of Moldova	TG	Togo
BB	Barbados	GH	Ghana	MG	Madagascar	TJ	Tajikistan
BE	Belgium	GN	Guinea	MK	The former Yugoslav Republic of Macedonia	TM	Turkmenistan
BF	Burkina Faso	GR	Greece	ML	Mali	TR	Turkey
BG	Bulgaria	HU	Hungary	MN	Mongolia	TT	Trinidad and Tobago
BJ	Benin	IE	Ireland	MR	Mauritania	UA	Ukraine
BR	Brazil	IL	Israel	MW	Malawi	UG	Uganda
BY	Belarus	IS	Iceland	MX	Mexico	US	United States of America
CA	Canada	IT	Italy	NE	Niger	UZ	Uzbekistan
CF	Central African Republic	JP	Japan	NL	Netherlands	VN	Viet Nam
CG	Congo	KE	Kenya	NO	Norway	YU	Yugoslavia
CH	Switzerland	KG	Kyrgyzstan	NZ	New Zealand	ZW	Zimbabwe
CJ	Côte d'Ivoire	KP	Democratic People's Republic of Korea	PL	Poland		
CM	Cameroon	KR	Republic of Korea	PT	Portugal		
CN	China	KZ	Kazakhstan	RO	Romania		
CU	Cuba	LC	Saint Lucia	RU	Russian Federation		
CZ	Czech Republic	LI	Liechtenstein	SD	Sudan		
DE	Germany	LK	Sri Lanka	SE	Sweden		
DK	Denmark	LR	Liberia	SG	Singapore		
EE	Estonia						

HAIR CARE COMPOSITIONS COMPRISING OPTICAL BRIGHTENERS AND NON-VOLATILE SOLVENTS

5

TECHNICAL FIELD

10 The present invention relates to a hair care composition comprising optical brighteners which alter the color of the hair, while enhancing the shininess of the hair, and protecting the hair from further damage. More specifically, the present invention relates to a hair care composition comprising optical brighteners and non-volatile solvents.

15

BACKGROUND

The desire to regain the natural color and shine of damaged hair and the desire to alter the color of the hair to be more appealing are widely held. Damaged hair is perceived by the consumer as unfavorable appearances and less manageability of the hair. Such unfavorable appearances include alteration and fading of original color, less shine, and less luster.

20

A common way for alleviating the unfavorable appearances of damaged hair and to achieve appealing hair color is to dye the hair to the color desired. Dyeing the hair would provide the consumer with a stable color of hair for a relatively long period. However, dyeing the hair is generally time-consuming, cumbersome, and messy. Dyestuff may also be chemically harsh to the hair, scalp, and skin. The hair can be further damaged by dyeing. Thus, hair dye products are not suitable for daily use. Further, dyeing can leave the hair with a dull appearance, making the hair look less shiny.

25

Based on the foregoing, there is a need for a hair composition which can be used daily and which can alter the color of the hair, while enhancing the shininess of the hair, and protecting the hair from further damage.

30

Use of optical brighteners, or compounds otherwise described by names such as fluorescent whitening agents, fluorescent brighteners, or fluorescent dyes, in the hair care field has been known in the art, such as in United States Patent 3,658,985, United States Patent 4,312,855, Canadian Patent 1,255,603,

35

United States Patent 3,577,528, Great Britain Patent Specification 1,328,108, South African Application 676,049, European Publication 87,060, and Great Britain Patent Specification 2,307,639.

40 However, the addition of large amounts of optical brighteners to hair compositions can actually lessen the hair's shininess if the amount of optical brightener is such that it crystallizes after the aqueous components of the composition have evaporated or the hair has dried. Such crystallization can make the hair look dull or "dusty". Accordingly, there is a need for a hair care composition comprising optical brighteners in which a high deposition of the
45 optical brightener on the hair is attainable, and yet in which a dulling or dusting of the hair's shininess is avoided.

None of the existing art provides all of the advantages and benefits of the present invention.

50

SUMMARY

The present invention is directed to a hair care composition comprising:
(a) an effective amount of an optical brightener; and (b) a non-volatile solvent for solubilizing at least 0.01 wt% of the optical brightener; wherein the optical brightener is deposited on the hair without crystallizing.

55

These and other features, aspects, and advantages of the present invention will become evident to those skilled in the art from a reading of the present disclosure.

DETAILED DESCRIPTION

60

While the specification concludes with claims particularly pointing and distinctly claiming the invention, it is believed that the present invention will be better understood from the following description.

65

All percentages are by weight of the total composition unless otherwise indicated. All ratios are weight ratios unless otherwise indicated. All percentages, ratios, and levels of ingredients referred to herein are based on the actual amount of the ingredient, and do not include solvents, fillers, or other materials with which the ingredient may be combined as commercially available products, unless otherwise indicated.

As used herein, "comprising" means that other steps and other ingredients
70 which do not affect the end result can be added. This term encompasses the
terms "consisting of" and "consisting essentially of".

All cited references are incorporated herein by reference in their entireties.
Citation of any reference is not an admission regarding any determination as to
its availability as prior art to the claimed invention.

75 OPTICAL BRIGHTENERS

Optical brighteners are compounds which absorb ultraviolet light and re-
emit the energy in the form of visible light. Specifically, the optical brighteners
useful herein have an absorption, preferably a major absorption peak, between a
wavelength of about 1nm and about 420nm, and an emission, preferably a major
80 emission peak, between a wavelength of about 360nm and about 830nm;
wherein the major absorption peak has a shorter wavelength than the major
emission peak. More preferably, the optical brighteners useful herein have a
major absorption peak between a wavelength of about 200nm and about 420nm,
and a major emission peak between a wavelength of about 400nm and about
85 780nm. Optical brighteners may or may not have a secondary absorption peak
in the visible range between a wavelength of about 360nm and about 830nm.
Optical brighteners can be described by other names in the art and in other
industries, such as fluorescent whitening agents, fluorescent brighteners, and
fluorescent dyes.

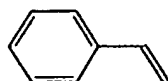
90 When applied to hair via suitable vehicles, optical brighteners herein
provide benefits to the hair in three areas. First, optical brighteners herein alter
the color of the hair by emitting light in the visible range. Second, optical
brighteners herein enhance the shine of the hair by emitting light in the visible
range. Third, optical brighteners herein protect the hair from ultraviolet light by
95 absorbing ultraviolet light.

Optical brighteners in general are based on the structures of aromatic and
heteroaromatic systems which provide these unique characteristics. The optical
brighteners useful in the present invention can be classified according to their
base structures, as described hereafter. Preferable optical brighteners herein
100 include polystyrylstilbenes, triazinstilbenes, hydroxycoumarins, aminocoumarins,
triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, and imidazoles.

Preferably, optical brighteners herein are included in the hair care composition of the present invention at a level by weight of from about 0.001% to about 20%, more preferably from about 0.01% to about 10%.

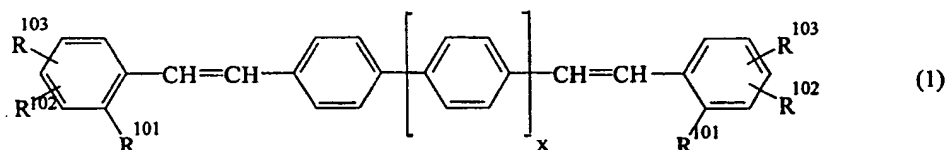
105 Polystyrylstilbenes

Polystyrylstilbenes are a class of compounds having two or more of the following base structure:



110

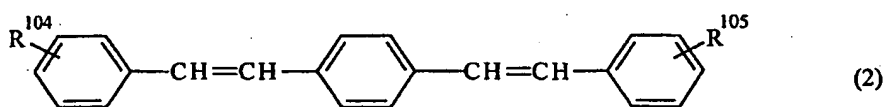
Polystyrylstilbenes useful in the present invention include those having formulae (1), (2) and (3):



115

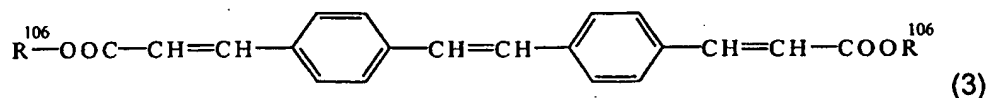
wherein R^{101} is H, OH, SO_3M , COOM, OSO_3M , $OPO(OH)OM$, wherein M is H, Na, K, Ca, Mg, ammonium, mono-, di-, tri- or tetra- C_1 - C_{30} -alkylammonium, mono-, di- or tri- C_1 - C_{30} -hydroxyalkylammonium or ammonium that is di- or tri-substituted with by a mixture of C_1 - C_{30} -alkyl and C_1 - C_{30} -hydroxyalkyl groups; or $SO_2N(C_1$ - C_{30} -alkyl) $_2$, O -(C_1 - C_{30} -alkyl), CN, Cl, $COO(C_1$ - C_{30} -alkyl), $CON(C_1$ - C_{30} -alkyl) $_2$ or $O(CH_2)_3N^+(CH_3)_2X^-$ wherein X^- is an anion of a chloride, bromide, iodide, formate, acetate, propionate, glycolate, lactate, acrylate, methanephosphonate, phosphite, dimethyl or diethyl phosphite anion; CN, or alkyl of 1 to 30 carbons, R^{102} and R^{103} , independently, are H, SO_3M wherein M is as previously defined; and x is 0 or 1; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably x is 1, R^{101} is SO_3Na and R^{102} and R^{103} are H; wherein the compound has a trans-coplanar orientation;

125



130

wherein R¹⁰⁴ and R¹⁰⁵, independently, are CN, COO(C₁-C₃₀-alkyl), CONHC₁-C₄-alkyl, or CON(C₁-C₄-alkyl)₂, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R¹⁰⁴ and R¹⁰⁵ is 2-cyano,
 135 wherein the compound has a trans-coplanar orientation; and



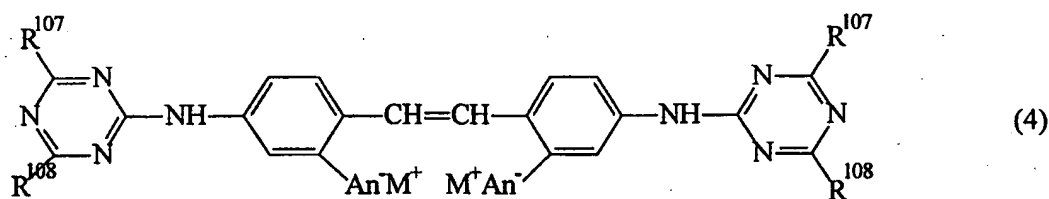
wherein each R¹⁰⁶, independently, is H, or alkyl of 1 to 30 carbons; and wherein
 140 the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation.

Suitable polystyrylstilbenes include disodium-1, 4'-bis(2-sulfostryl) bisphenyl (C.I. Fluorescent Brightener 351) with tradename Tinopal CBS-X available from Ciba Specialty Chemicals, 1,4-bis(2-cyanostyryl)benzene (C.I.
 145 Fluorescent Brightener 199), with tradename Ultraphor RN available from BASF.

Triazinstilbenes

Triazinstilbenes are a class of compounds having both triazin and stilbene structures in the same molecule.

Triazinstilbenes useful in the present invention include those having
 150 formulae (4):



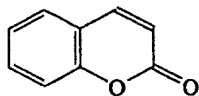
wherein R¹⁰⁷ and R¹⁰⁸, independently, are phenylamino, mono- or disulfonated
 155 phenylamino, morpholino, N(CH₂CH₂OH)₂, N(CH₃)(CH₂CH₂OH), NH₂, N(C₁-C₄-alkyl)₂, OCH₃, Cl, NH-(CH₂)₁₋₄SO₃H or NH-(CH₂)₁₋₄OH; An⁻ is an anion of a carboxylate, sulfate, sulfonate, or phosphate, and M is as previously defined, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R¹⁰⁷ is 2, 5-disulfophenylamino and each R¹⁰⁸ is
 160 morpholino; or each R¹⁰⁷ is 2, 5-disulfophenylamino and each R¹⁰⁸ is N(C₂H₅)₂; or each R¹⁰⁷ is 3-sulfophenyl and each R¹⁰⁸ is NH(CH₂CH₂OH) or

$N(CH_2CH_2OH)_2$; or each R^{107} is 4-sulfophenyl and each R^{108} is $N(CH_2CH_2OH)_2$; and in each case, the sulfo group is SO_3M in which M is sodium; wherein the compound has a trans-coplanar orientation.

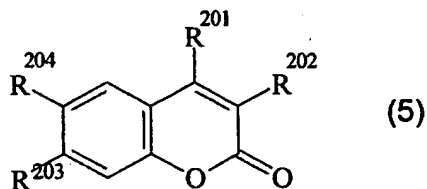
165 Suitable triazin stilbenes include 4,4'-bis-[(4-anilino-6-bis(2-hydroxyethyl)amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acid with tradename Tinopal UNPA-GX available from Ciba Specialty Chemicals, 4,4'-bis-[(4-anilino-6-morpholine-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate with tradename Tinopal AMS-GX available from Ciba Specialty Chemicals, 4,4'-bis-[(4-anilino-6-(2-hydroxyethyl)methyl amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate with tradename Tinopal 5BM-GX available from Ciba Specialty Chemicals, 4,4'-bis-[(4,6-dianilino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate, 4,4'-bis-[(4-anilino-6-methylamino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate, 4,4'-bis-[(4-anilino-6-ethylamino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disodium sulfonate, and 4,4'-bis(4-phenyl-1,2,3-triazol-2-yl)stilbene-2,2'-disulfonic acid.

Hydroxycoumarins

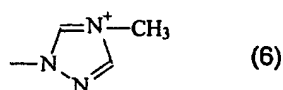
Hydroxycoumarins are a class of compounds having the following base coumarin structure and having at least one hydroxy moiety:



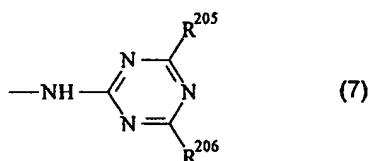
Hydroxycoumarins useful in the present invention include those having formulae (5):



wherein R^{201} is H, OH, Cl, CH_3 , CH_2COOH , CH_2SO_3H , CH_2OSO_3H , or $CH_2OPO(OH)OH$, R^{202} is H, phenyl, $COO-C_1-C_{30}$ -alkyl, glucose, or a group of formula (6):



and R²⁰³ is OH, or O-C₁-C₃₀-alkyl, and R²⁰⁴ is OH or O-C₁-C₃₀ alkyl,
 195 glycoside, or a group of the following formula (7):



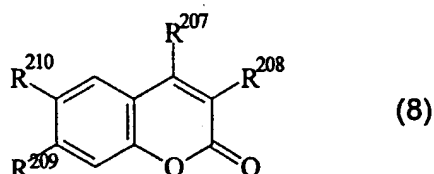
wherein R²⁰⁵ and R²⁰⁶ are independently, phenylamino, mono- or disulfonated
 200 phenylamino, morpholino, N(CH₂CH₂OH)₂, N(CH₃)(CH₂CH₂OH), NH₂, N(C₁-
 C₃₀-alkyl)₂, OCH₃, Cl, NH-(CH₂)₁₋₄SO₃H or NH-(CH₂)₁₋₄OH.

Suitable hydroxycoumarins include 6,7-dihydroxycoumarin available from
 Wako Chemicals, 4-methyl-7-hydroxycoumarin available from Wako Chemicals,
 4-methyl-6,7-dihydroxycoumarin available from Wako Chemicals, esculin
 205 available from Wako Chemicals, and umbelliferone (4-hydroxycoumarin)
 available from Wako Chemicals.

Aminocoumarins

Aminocoumarins are a class of compounds having the base coumarin
 structure and having at least one amino moiety.

210 Aminocoumarins useful in the present inventions include those having
 formulae (8):

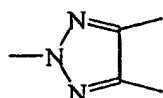


215 wherein R²⁰⁷ is H, Cl, CH₃, CH₂COOH, CH₂SO₃H, CH₂OSO₃H, or
 CH₂OPO(OH)OH, R²⁰⁸ is H, phenyl, or COOC₁-C₃₀ alkyl, and R²⁰⁹ and R²¹⁰
 are independently H, NH₂, N(C₁-C₃₀alkyl)₂, NHC₁-C₃₀alkyl, or NHCOC₁-
 C₃₀alkyl.

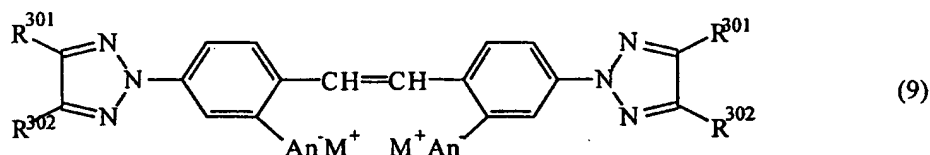
220 Suitable aminocoumarins include 4-methyl-7,7'-diethylamino coumarin with tradename Calcofluor-RWP available from BASF, 4-methyl-7,7'-dimethylamino coumarin with tradename Calcofluor-LD available from BASF.

Triazoles

Triazoles are a class of compounds having the following base structure:

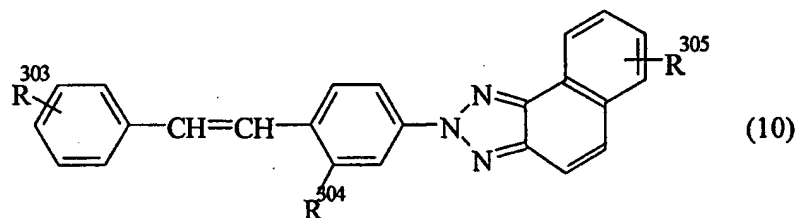


Triazoles useful in the present inventions include those having formulae (9) through (12) and (15) through (20):



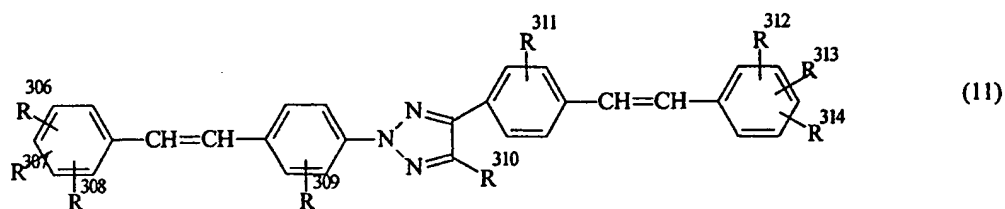
wherein R³⁰¹ and R³⁰², independently, are H, C₁-C₃₀alkyl, phenyl or monosulfonated phenyl; An⁻ and M are as previously defined, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R³⁰¹ is phenyl, R³⁰² is H and M is sodium; wherein the compound has a trans-coplanar orientation;

235



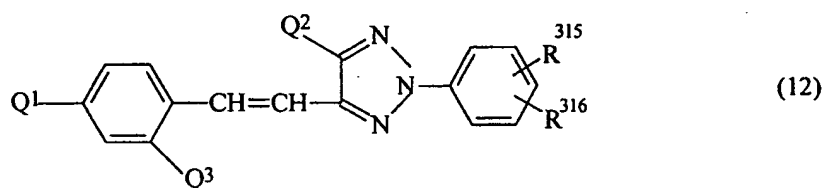
240 wherein R³⁰³ is H or Cl; R³⁰⁴ is SO₃M, SO₂N(C₁-C₃₀alkyl)₂, SO₂O-phenyl or CN; R³⁰⁵ is H, SO₃M, COOM, OSO₃M, or OPO(OH)OM; and M is as previously defined, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R³⁰³ and R³⁰⁵ are H and R³⁰⁴ is SO₃M in which M is Na; wherein the compound has a trans-coplanar orientation;

245

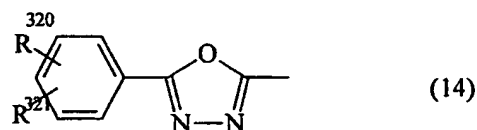
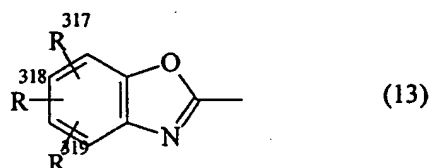


wherein each of R³⁰⁶ and R³¹² independently represents H, a sulfonic acid group or the salts, esters or amides thereof, a carboxylic acid group or the salts, esters or amides thereof, a cyano group, a halogen atom, an unsubstituted or substituted alkylsulfonyl, arylsulfonyl, alkyl, alkoxy, aralkyl, aryl, aryloxy, aralkoxy or cycloalkyl radical, an unsubstituted or substituted 5-membered heterocyclic ring containing 2 to 3 nitrogen atoms or one oxygen atom and 1 or 2 nitrogen atoms, or together with R³⁰⁷ and R³¹³ they represent a methylenedioxy, ethylenedioxy, methylenoxymethylenoxy, trimethylene, tetramethylene, propenylene, butenylene or butadienylene radical, each of R³⁰⁷ and R³¹³ independently represents H, a sulfonic acid group or the salts, esters or amides thereof, a carboxylic acid group or the salts, esters or amides thereof, a cyano group, a halogen atom, an unsubstituted or substituted alkyl or alkoxy radical, or together with R³⁰⁶ and R³¹² represent a methylenedioxy, ethylenedioxy, methylenoxymethylenoxy, trimethylene, tetramethylene, propenylene, butenylene or butadienylene radical, each of R³⁰⁸ and R³¹⁴ independently represents H, a halogen atom or an unsubstituted or substituted alkyl radical, each of R³⁰⁹ and R³¹¹ independently represents H, a halogen atom, a cyano group a sulfonic acid group or the salts, esters or amides thereof, or a carboxylic acid group or the salts, esters or amides thereof, and R³¹⁰ independently represents H, a halogen atom, a cyano group a sulfonic acid group or the salts, alkyl radicals preferably by hydroxy, alkoxy of 1 to 30 carbon atoms, cyano, halogen, carboxy, sulfonic acid groups, carbalkoxy having 1 to 30 carbon atoms in the alkoxy moiety, phenyl or phenoxy; alkoxy radicals can be substituted by hydroxy, alkoxy of 1 to 30 carbon atoms, cyano, halogen, carboxy, carbalkoxy having 1 to 30 carbon atoms in the alkoxy moiety, phenyl or phenoxy; phenyl, phenylalkyl or phenoxy radicals can be substituted by halogen, cyano, carboxy, carbalkoxy having 1 to 30 carbon atoms in the alkoxy moiety, sulfo, or alkyl or alkoxy each of 1 to 30 carbon atoms; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; possible cycloalkyl radicals are preferably cyclohexyl and cyclopentyl

radicals which can be substituted by alkyl of 1 to 30 carbon atoms; possible 5-membered heterocyclic rings are v-triazole, oxazole or 1, 3, 4- oxdiazole radicals which can contain as substituents alkyl radicals of 1 to 4 carbon atoms, halogen, phenyl, carboxy, carbalkoxy having 1 to 30 carbon atoms in the alkoxy moiety, cyano, benzyl, alkoxy of 1 to 30 carbon atoms, phenoxy or sulfo, whilst two adjacent substituents of the triazole and oxazole radicals together are able to form a substituted or unsubstituted fused benzene nucleus; wherein the compound has a trans-coplanar orientation;

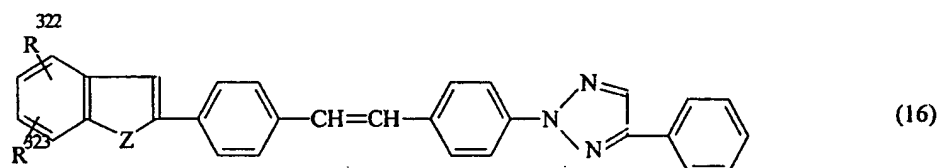
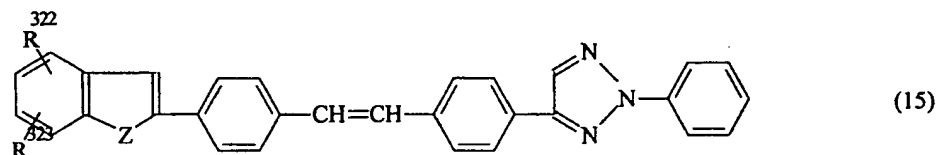


wherein Q¹ denotes one of the ring systems (13) or (14);

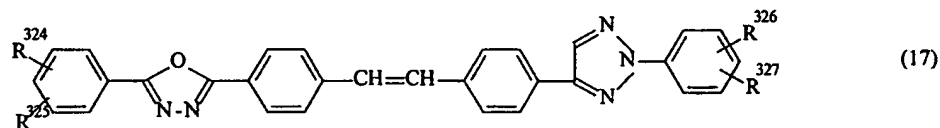


and wherein R³¹⁷ denotes H, alkyl with 1 to 30 carbon atoms, cyclohexyl, phenylalkyl with C₁-C₃₀ carbon atoms in the alkyl part, phenyl, alkoxy with 1 to 30 carbon atoms, or Cl, or, conjointly with R³¹⁸, denotes alkylene with 3 to 30 carbon atoms, R³¹⁸ denotes H or alkyl with 1 to 30 carbon atoms or, conjointly with R³¹⁷, denotes alkylene with 3 to 30 carbon atoms, R³¹⁹ denotes H or methyl, R³²⁰ denotes H, alkyl with 1 to 30 carbon atoms, phenyl, alkoxy with 1 to 30 carbon atoms, or Cl, or, conjointly with R³²¹, denotes a fused benzene ring, R³²¹ denotes H or Cl or conjointly with R³²⁰, denotes a fused benzene ring, R³¹⁵ denotes H, alkyl with 1 to 30 carbon atoms, alkoxy with 1 to 30 carbon atoms or Cl, R³¹⁶ denotes H or Cl, Q² denotes H, Cl alkyl with 1 to 30 carbon atoms or phenyl and Q³ denotes H or Cl; wherein the compound has a trans-

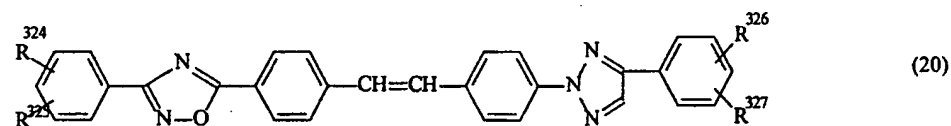
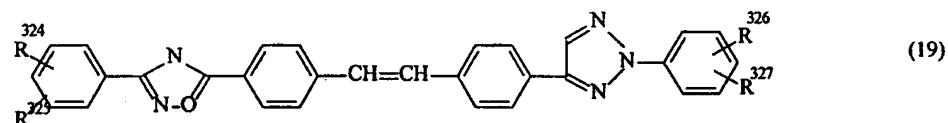
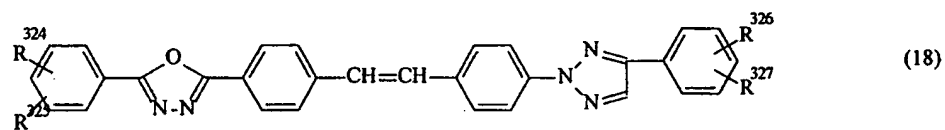
coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar
 305 orientation;



310 wherein R³²² denotes H, Cl, methyl, phenyl, benzyl, cyclohexyl or methoxy,
 R³²³ denotes H or methyl and Z denotes O or S; wherein the compound has a
 trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar
 orientation; and



315



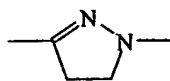
320 wherein R³²⁴ denotes H, Cl, alkyl with 1 to 30 carbon atoms, phenylalkyl with 1
 to 30 carbon atoms, phenyl or alkoxy with 1 to 30 carbon atoms, or R³²⁴
 conjointly with R³²⁵ denotes a fused benzene radical, R³²⁵ denotes H or methyl
 or R³²⁵ conjointly with R³²⁴ denotes a fused benzene radical, R³²⁶ denotes H,
 alkyl with 1 to 30 carbon atoms, alkoxy with 1 to 30 carbon atoms, Cl, carbalkoxy

325 with 1 to 30 carbon atoms or alkylsulfonyl with 1 to 30 carbon atoms and R^{327} denotes H, Cl, methyl or methoxy; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation.

Suitable triazoles include 2-(4-styryl-3-sulfophenyl)-2H-naptho[1,2-d]triazole (C.I. Fluorescent Brightener 46) with tradename Tinopal RBS available
330 from Ciba Specialty Chemicals.

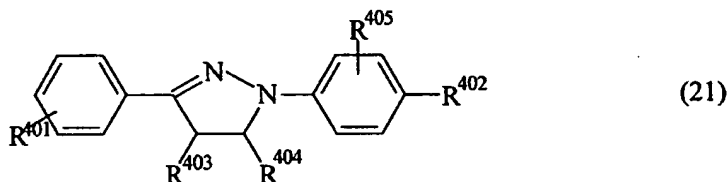
Pyrazolines

Pyrazolines are a class of compounds having the following base structure:



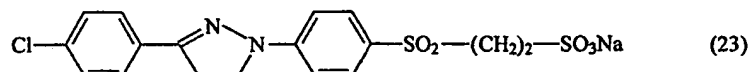
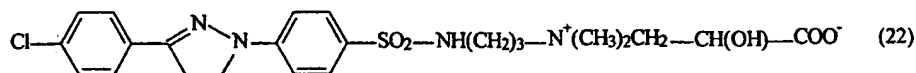
335

Pyrazolines useful in the present invention include those having formulae (21) through (23):



340

wherein R^{401} is H, Cl or $N(C_1-C_{30}\text{-alkyl})_2$, R^{402} is H, Cl, SO_3M , SO_2NH_2 , $SO_2NH-(C_1-C_{30}\text{alkyl})$, $COO-C_1-C_{30}\text{alkyl}$, $SO_2-C_1-C_{30}\text{alkyl}$, $SO_2NH(CH_2)_{1-4}N^+(CH_3)_3$ or $SO_2(CH_2)_{1-4}N^+H(C_1-C_{30}\text{-alkyl})_2An^-$, R^{403} and R^{404} are the same or different and each is H, $C_1-C_{30}\text{alkyl}$ or phenyl and R^{405} is H or Cl; and
345 An^- and M are as previously defined, preferably R^{401} is Cl, R^{402} is $SO_2CH_2CH_2N^+H(C_1-C_4\text{-alkyl})_2An^-$ in which An^- is phosphite and R^{403} , R^{404} and R^{405} are each H; and formulae (22) and (23) shown below.



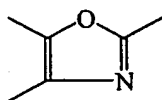
350

Suitable pyrazolines include 1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline (C.I. Fluorescent Brightener 121) with tradename Blankophor DCB

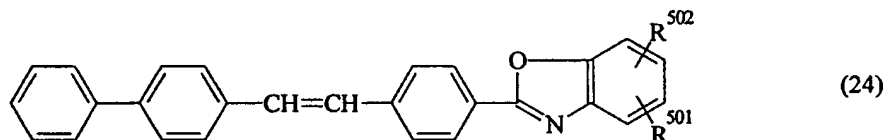
available from Bayer, 1-[4-(2-sulfoethylsulfonyl)phenyl]-3-(4-chlorophenyl)-2-pyrazoline, 1-[4-(2-sulfoethylsulfonyl)phenyl]-3-(3,4-dichloro-6-methylphenyl)-2-pyrazoline, 1-<4-{N-[3-(N,N,N-trimethylammonio)propyl]-amidosulfonyl}phenyl>-3-(4-chlorophenyl)-2-pyrazoline methylsulfate, and 1-<4-{2-[1-methyl-2-(N,N-dimethylamino)ethoxy]ethylsulfonyl}phenyl>-3-(4-chlorophenyl)-2-pyrazoline methylsulfate.

360 Oxazoles

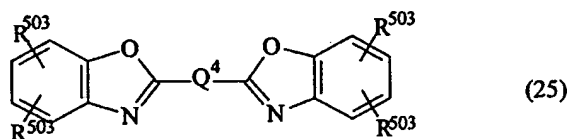
Oxazoles are a class of compounds having the following base structure:



365 Oxazoles useful in the present inventions include those having formulae (24), (25), (26) and (27):

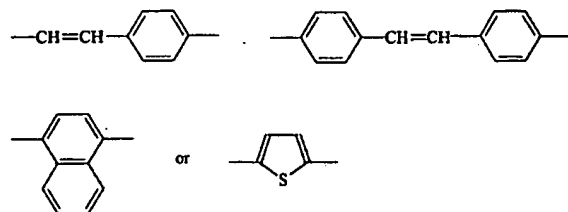


370 wherein R⁵⁰¹ and R⁵⁰², independently, are H, Cl, C₁-C₃₀alkyl or SO₂-C₁-C₃₀alkyl, wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R⁵⁰¹ is 4-CH₃ and R⁵⁰² is 2-CH₃ wherein the compound has a trans-coplanar orientation;

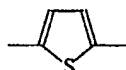


375 wherein R⁵⁰³, independently, is H, C(CH₃)₃, C(CH₃)₂-phenyl, C₁-C₃₀alkyl or COO-C₁-C₃₀alkyl, preferably H and Q⁴ is -CH=CH-;

380

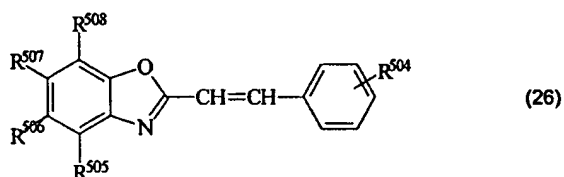


preferably



385 or one group R⁵⁰³ in each ring is 2-methyl and the other R⁵⁰³ is H and Q⁴ is ---CH=CH--- ; or one group R⁵⁰³ in each ring is 2-C(CH₃)₃ and the other R⁵⁰³ is H; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation;

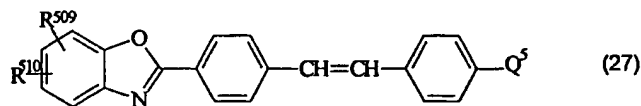
390



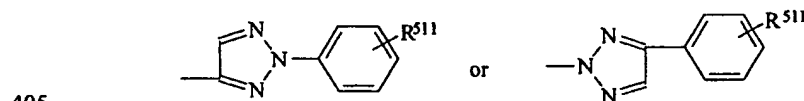
395

wherein R⁵⁰⁴ is CN, Cl, COO-C₁-C₃₀alkyl or phenyl; R⁵⁰⁵ and R⁵⁰⁶ are the atoms required to form a fused benzene ring or R⁵⁰⁶ and R⁵⁰⁸, independently, are H or C₁-C₃₀alkyl; and R⁵⁰⁷ is H, C₁-C₃₀alkyl or phenyl; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation; preferably R⁵⁰⁴ is a 4-phenyl group and each of R⁵⁰⁵ to R⁵⁰⁸ is H; wherein the compound has a trans-coplanar orientation; and

400



wherein R⁵⁰⁹ denotes H, Cl, alkyl with 1 to 30 carbon atoms, cyclohexyl, phenylalkyl with 1 to 3 carbon atoms in the alkyl part, phenyl or alkoxy with 1 to 30 carbon atoms, R⁵¹⁰ denotes H or alkyl with 1 to 30 carbon atoms, and Q⁵ denotes a radical;

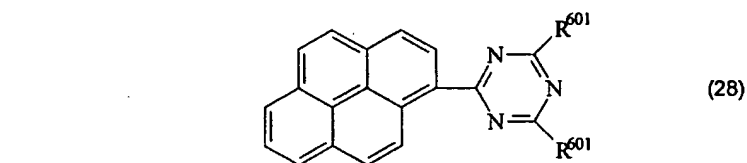


wherein R⁵¹¹ represents H, alkyl with 1 to 30 carbon atoms, alkoxy with 1 to 30 carbon atoms, Cl, carbalkoxy with 1 to 30 carbon atoms, unsubstituted sulfamoyl or sulfamoyl which is monosubstituted or disubstituted by alkyl or hydroxyalkyl with 1 to 30 carbon atoms or represents alkylsulfonyl with 1 to 30 carbon atoms; wherein the compound has a trans-coplanar orientation or cis-coplanar orientation, preferably a trans-coplanar orientation.

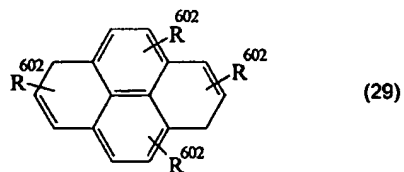
Suitable oxazoles include 4,4'-bis(5-methylbenzoxazol-2-yl)stilbene, and 2-(4-methoxycarbonylstyryl)benzoxazole.

415 Pyrenes

Pyrenes useful in the present invention include those having formulae (28) and (29):



wherein each R⁶⁰¹, independently, is C₁-C₃₀alkoxy; preferably methoxy; and



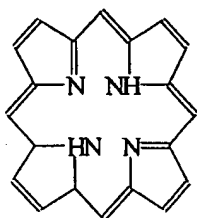
425 wherein each R⁶⁰², independently, is H, OH, or SO₃M, wherein M is as previously defined, sulfonated phenylamino, or anilino.

Suitable pyrenes include 2,4-dimethoxy-6-(1'-pyrenyl)-1,3,5-triazine (C.I. Fluorescent Brightener 179) with tradename Fluolite XMF, 8-hydroxy-1,3,6-pyrenetrisulfonic acid (D&C Green No.8), and 3-hydroxy-5,8,10-trisulphanilic pyrene.

430

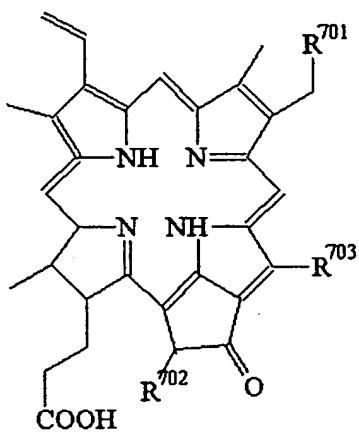
Porphyrins

Porphyrins useful in the present invention include those having formulae (30), (31), and (32):



(30)

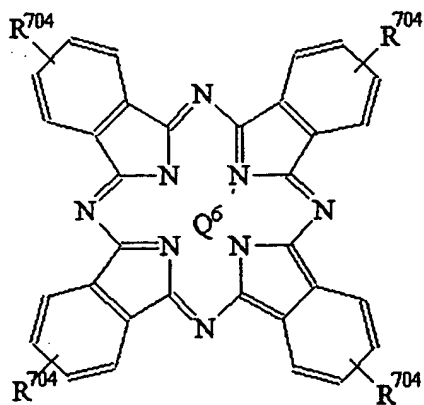
435



(31)

440

wherein R^{701} is CH_3 or CHO , R^{702} is H or $COOC_1-C_{30}$ alkyl, and R^{703} is H or an alkyl group having 1 to 30 carbons; and



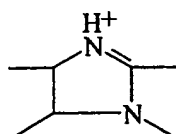
(32)

wherein each R^{704} , independently, is H, SO_3M , $COOM$, OSO_3M , or
 445 $OPO(OH)OM$, wherein M is as previously defined, halide, or alkyl of 1 to 30
 carbons; and Q^6 is Cu, Mg, Fe, Cr, Co, or mixtures thereof with cationic charges.

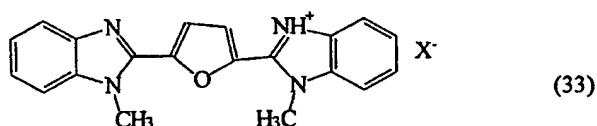
Suitable porphyrins include porphyrin available from Wako Chemicals and
 Copper II phthalocyanine available from Wako Chemicals.

Imidazoles

450 Imidazoles are a class of compounds having the following base structure:



455 Imidazoles useful in the present invention include those having formulae
 (33):



460 wherein X^- is as previously defined.

Suitable imidazoles include those with tradename of C.I. Fluorescence
 Brightener 352, or Uvtext AT available from Ciba Speciality Chemical.

NON-VOLATILE SOLVENTS

The non-volatile solvents solubilize the optical brightener. As used herein,
 465 "non-volatile" means having a boiling point greater than about $100^\circ C$. Preferably
 the non-volatile solvent is capable of dissolving at least 0.01 wt% of the optical
 brightener.

Optical brighteners in a water or ethanol solution, without a non-volatile
 solvent, can make the hair look dull or dusty after the water or ethanol has
 470 evaporated, i.e., after the hair has dried. This dullness or dustiness results from
 crystallization of the optical brightener on the hair. If a non-volatile solvent is not
 added to the composition, the level of optical brightener in the composition must
 be reduced in order to prevent crystallization and the consequent dulling or
 dusting of the hair. This, however, is undesirable, because reduction in the level

475 of optical brightener causes a reduction in the benefits (e.g., color enhancement, shine, UV protection) provided by the optical brightener.

Without being bound by theory, it is believed that the addition of a nonvolatile solvent as described herein reduces the crystallization effect, and thus permits a high level of the optical brightener to be added to the hair care
480 composition, and thus deposited on the hair, but without compromising the benefits provided by the optical brightener.

Non-volatile solvents useful herein include alkyl alcohols having more than 3 carbons, and polyhydric alcohols. The polyhydric alcohols useful herein include 1,2-propane diol or propylene glycol, 1,3-propane diol, hexylene glycol, glycerin, diethylene glycol, dipropylene glycol, 1,2-butylene glycol, and 1,4-
485 butylene glycol. Preferably the hair care compositions of the present invention contain at least about 0.1 wt% of the non-volatile solvent.

CARRIER

The hair care compositions of the present invention may comprise a carrier.
490 The level and species of the carrier are selected according to the compatibility with other components, and desired characteristic of the product.

The carrier useful in the present invention include volatile solvents, propellants, and mixtures thereof.

Volatile solvents useful herein include water, lower alkyl alcohols having
495 from 1 to 3 carbons, and hydrocarbons having from about 5 to about 8 carbons. The preferred volatile solvents are water, ethanol, isopropanol, pentane, hexane, and heptane. The water useful herein include deionized water and water from natural sources containing mineral cations. Deionized water is preferred.

Propellants may be used for mousse and hair spray product forms.
500 Propellants, when used in the present invention, are selected depending on variables such as the remainder of components, the package, and whether the product is designed to be used standing or invert.

Propellants useful herein include fluorohydrocarbons such as difluoroethane 152a (supplied by DuPont), dimethylether, and hydrocarbons such
505 as propane, isobutane, n-butane, mixtures of hydrocarbons such as LPG (liquefied petroleum gas), carbon dioxide, nitrous oxide, nitrogen, and compressed air.

COMPOSITIONS

The hair care compositions of the present invention may comprise a component which characterizes the form of the product. Product forms useful herein include, but are not limited to, shampoo, conditioner, treatment, mousse, spray, lotion, gel, and cream products, all of which can be designed for rinse-off or leave-on convenience. Product functions useful herein include, but are not limited to, cleansing, and conditioning products. For example, a deterative surfactant, a hair conditioning agent, and mixtures thereof may be comprised. The components and their levels are selected by one skilled in the art depending on the desired characteristic of the product.

DETERGENT SURFACTANT

Compositions herein may include a detergent surfactant. The detergent surfactants herein are those suitable for cleansing the hair. Detergent surfactants useful herein include anionic surfactants, amphoteric and zwitterionic surfactants, and nonionic surfactants.

The detergent surfactants, when present, are preferably included at a level of from about 0.01% to about 75% by weight of the composition. Two or more surfactants can be used.

Anionic Surfactant

Anionic surfactants useful herein include alkyl and alkyl ether sulfates. These materials have the respective formulae ROSO_3M and $\text{RO}(\text{C}_2\text{H}_4\text{O})_x\text{SO}_3\text{M}$, wherein R is alkyl or alkenyl of from about 8 to about 30 carbon atoms, x is 1 to about 10, and M is hydrogen or a cation such as ammonium, alkanolammonium (e.g., triethanolammonium), a monovalent metal cation (e.g., sodium and potassium), or a polyvalent metal cation (e.g., magnesium and calcium). Preferably, M should be chosen such that the anionic surfactant component is water soluble. The anionic surfactant or surfactants should be chosen such that the Krafft temperature is about 15°C or less, preferably about 10°C or less, and more preferably about 0°C or less. It is also preferred that the anionic surfactant be soluble in the composition hereof.

Krafft temperature refers to the point at which solubility of an ionic surfactant becomes determined by crystal lattice energy and heat of hydration, and corresponds to a point at which solubility undergoes a sharp, discontinuous increase with increasing temperature. Each type of surfactant will have its own characteristic Krafft temperature. Krafft temperature for ionic surfactants is, in

general, well known and understood in the art. See, for example, Myers, Drew, Surfactant Science and Technology, pp. 82-85, VCH Publishers, Inc. (New York, New York, USA), 1988 (ISBN 0-89573-399-0), which is incorporated by reference
545 herein in its entirety.

In the alkyl and alkyl ether sulfates described above, preferably R has from about 8 to about 18 carbon atoms in both the alkyl and alkyl ether sulfates. The alkyl ether sulfates are typically made as condensation products of ethylene
550 oxide and monohydric alcohols having from about 8 to about 24 carbon atoms. The alcohols can be derived from fats, e.g., coconut oil, palm oil, tallow, or the like, or the alcohols can be synthetic. Lauryl alcohol and straight chain alcohols derived from coconut oil and palm oil are preferred herein. Such alcohols are reacted with 1 to about 10, and especially about 3, molar proportions of ethylene
555 oxide and the resulting mixture of molecular species having, for example, an average of 3 moles of ethylene oxide per mole of alcohol, is sulfated and neutralized.

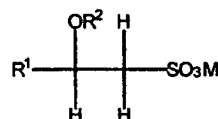
Specific examples of alkyl ether sulfates which can be used are sodium and ammonium salts of coconut alkyl triethylene glycol ether sulfate; tallow alkyl
560 triethylene glycol ether sulfate, and tallow alkyl hexaoxyethylene sulfate. Highly preferred alkyl ether sulfates are those comprising a mixture of individual compounds, said mixture having an average alkyl chain length of from about 8 to about 16 carbon atoms and an average degree of ethoxylation of from 1 to about 4 moles of ethylene oxide. Such a mixture also comprises from 0% to about 20%
565 by weight C₁₂₋₁₃ compounds; from about 60% to about 100% by weight of C₁₄₋₁₅₋₁₆ compounds, from 0% to about 20% by weight of C₁₇₋₁₈₋₁₉ compounds; from about 3% to about 30% by weight of compounds having a degree of ethoxylation of 0; from about 45% to about 90% by weight of compounds having a degree of ethoxylation of from 1 to about 4; from about 10%
570 to about 25% by weight of compounds having a degree of ethoxylation of from about 4 to about 8; and from about 0.1% to about 15% by weight of compounds having a degree of ethoxylation greater than about 8.

Other suitable anionic surfactants are the water-soluble salts of organic, sulfuric acid reaction products of the general formula [R¹-SO₃-M] where R¹ is
575 selected from the group consisting of a straight or branched chain, saturated aliphatic hydrocarbon radical having from about 8 to about 24, preferably about 8 to about 18, carbon atoms; and M is as previously described above in this

section. Examples of such surfactants are the salts of an organic sulfuric acid reaction product of a hydrocarbon of the methane series, including iso-, neo-,
 580 and n-paraffins, having about 8 to about 24 carbon atoms, preferably about 8 to about 18 carbon atoms and a sulfonating agent, e.g., SO₃, H₂SO₄, obtained according to known sulfonation methods, including bleaching and hydrolysis. Preferred are alkali metal and ammonium sulfonated C₈₋₁₈ n-paraffins.

Still other suitable anionic surfactants are the reaction products of fatty
 585 acids esterified with isethionic acid and neutralized with sodium hydroxide where, for example, the fatty acids are derived from coconut or palm oil; or sodium or potassium salts of fatty acid amides of methyl tauride in which the fatty acids, for example, are derived from coconut oil. Other similar anionic surfactants are described in U.S. Patents 2,486,921, 2,486,922, and 2,396,278, which are
 590 incorporated by reference herein in their entirety.

Another class of anionic surfactants suitable for use in the shampoo compositions are the β-alkyloxy alkane sulfonates. These compounds have the following formula:

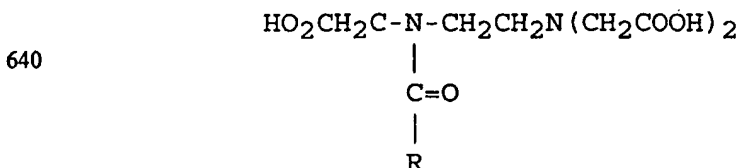


595 where R¹ is a straight chain alkyl group having from about 6 to about 20 carbon atoms, R² is a lower alkyl group having from about 1, preferred, to about 3 carbon atoms, and M is as hereinbefore described. Many other anionic surfactants suitable for use in the shampoo compositions are described in
 600 McCutcheon's, Emulsifiers and Detergents, 1989 Annual, published by M. C. Publishing Co., and in U.S. Patent 3,929,678, which descriptions are incorporated herein by reference in their entirety. Preferred anionic surfactants for use in the shampoo compositions include ammonium lauryl sulfate, ammonium laureth sulfate, triethylamine lauryl sulfate, triethylamine laureth sulfate,
 605 sulfate, triethanolamine lauryl sulfate, triethanolamine laureth sulfate, monoethanolamine lauryl sulfate, monoethanolamine laureth sulfate, diethanolamine lauryl sulfate, diethanolamine laureth sulfate, lauric monoglyceride sodium sulfate, sodium lauryl sulfate, sodium laureth sulfate, potassium lauryl sulfate, potassium laureth sulfate, ammonium cocoyl sulfate,
 610 ammonium lauroyl sulfate, sodium cocoyl sulfate, sodium lauroyl sulfate,

potassium cocoyl sulfate, potassium lauryl sulfate, triethanolamine lauryl sulfate, triethanolamine lauryl sulfate, monoethanolamine cocoyl sulfate, monoethanolamine lauryl sulfate, sodium tridecyl benzene sulfonate, and sodium dodecyl benzene sulfonate, and mixtures thereof.

615 Other anionic surfactants for use herein include polyhydrophilic anionic surfactants. By "polyhydrophilic" herein, is meant a surfactant that has at least two hydrophilic groups which provide a hydrophilic nature. Polyhydrophilic surfactants useful herein are only those having at least two hydrophilic groups in the molecule, and is not intended to encompass those which only have one
620 hydrophilic group. One molecule of the polyhydrophilic anionic surfactant herein may comprise the same hydrophilic groups, or different hydrophilic groups. Specifically, the polyhydrophilic anionic surfactants comprise at least one group selected from the group consisting of carboxy, hydroxy, sulfate, sulfonate, and phosphate. Suitable polyhydrophilic anionic surfactants are those which
625 comprise at least one of a carboxy, sulfate, or sulfonate group, more preferably those which comprise at least one carboxy group.

Nonlimiting examples of polyhydrophilic anionic surfactants include N-acyl-L-glutamates such as N-cocoyl-L-glutamate and, N-lauroyl-L-glutamate, laurimino dipropionate, N-acyl-L-aspartate, di-(N-lauroyl N-methyl taurate),
630 polyoxyethylene laurylsulfosuccinate, disodium N-octadecylsulfosuccinate; disodium lauryl sulfosuccinate; diammonium lauryl sulfosuccinate; tetra sodium N-(1,2-dicarboxyethyl)-N-octadecylsulfosuccinate; the diamyl ester of sodium sulfosuccinic acid; the dihexyl ester of sodium sulfosuccinic acid; and the dioctyl ester of sodium sulfosuccinic acid, and 2-cocoalkyl N-carboxyethyl N-
635 carboxyethoxyethyl imidazolinium betaine, lauroamphohydroxypropylsulfonate, cocoglyceryl ether salts, cocoglyceride sulfate, lauroyl isethionate, lauroamphoacetate, and those of the following formula:



645 wherein R is an alkyl of 8 to 18 carbons. Other polyhydrophilic anionic surfactants include olefin sulfonates having about 10 to about 24 carbon atoms. The term "olefin sulfonates" is used herein to mean compounds which can be produced by the sulfonation of alpha-olefins by means of uncomplexed sulfur trioxide, followed by neutralization of the acid reaction mixture in conditions such

650 that any sulfones which have been formed in the reaction are hydrolyzed to give the corresponding hydroxy-alkanesulfonates. The sulfur trioxide can be liquid or gaseous, and is usually, but not necessarily, diluted by inert diluents, for example by liquid SO₂, chlorinated hydrocarbons, etc., when used in the liquid form, or by air, nitrogen, gaseous SO₂, etc., when used in the gaseous form. The α-olefins

655 from which the olefin sulfonates are derived are mono-olefins having about 8 to about 24 carbon atoms, preferably about 10 to about 16 carbon atoms. Preferably, they are straight chain olefins. In addition to the true alkene sulfonates and a proportion of hydroxy-alkanesulfonates, the olefin sulfonates can contain minor amounts of other materials, such as alkene disulfonates

660 depending upon the reaction conditions, proportion of reactants, the nature of the starting olefins and impurities in the olefin stock and side reactions during the sulfonation process. A specific α-olefin sulfonate mixture of the above type is described more fully in U.S. Patent 3,332,880, to Pflaumer and Kessler, issued July 25, 1967, which is incorporated by reference herein in its entirety.

665 Another class of polyhydrophilic anionic surfactants are amino acid surfactants which are surfactants that have the basic chemical structure of an amino acid compound, i.e., that contains a structural component of one of the naturally-occurring amino acids. It is understood by the artisan that some surfactants may be regarded as both a polyhydrophilic anionic surfactant, and an

670 amino acid surfactant. These surfactants are suitable anionic surfactants.

Nonlimiting examples of amino acid surfactants include, N-cocoylalaninate, N-acyl-N-methyl-β-alanate, N-acylsarcosinate; N-alkylamino propionates and N-alkyliminodipropionates, specific examples of which include N-lauryl-β-amino propionic acid or salts thereof, and N-lauryl-β-imino-

675 dipropionate, N-acyl-DL-alaninate, sodium lauryl sarcosinate, sodium lauroyl
sarcosinate, lauryl sarcosine, cocoyl sarcosine, N-acyl-N-methyl taurate, lauroyl
taurate, and lauroyl lactylate.

Commercially available anionic surfactants suitable are N-acyl-L-
glutamate with a tradename AMISOFT CT-12S, N-acyl potassiumglycine with a
680 tradename AMILITE GCK-12, lauroyl glutamate with a tradename AMISOFT LS-
11, and N-acyl-DL-alaninate with tradename AMILITE ACT12 supplied by
Ajinomoto; acylaspartate with tradenames ASPARACK and AAS supplied by
Mitsubishi Chemical; and acyl derivatives of tradename ED3A supplied by
Hampshire Chemical Corp.

685 Optionally the counter ion of anionic surfactants may be polyvalent
cations. It has been found that these anionic surfactants, along with the cationic
conditioning agents, and polyvalent metal cations as described later, form a
coacervate in the compositions. Cationic conditioning agents may be included in
the present composition to provide a shampoo which both cleanse and condition
690 the hair from a single product.

Coacervate formulation is dependent upon a variety of criteria such as
molecular weight, component concentration, and ratio of interacting ionic
components ionic strength, charge density of the cationic and anionic
components, pH, and temperature. Coacervate systems and the effect of these
695 parameters are known in the art.

It is believed to be particularly advantageous, for the anionic surfactants
and the polyvalent metal cations at certain levels to be present with the cationic
conditioning agents in a coacervate phase. The coacervates formed in the
compositions are believed to readily deposit on the hair upon diluting the
700 coacervate with abundant water, i.e., rinsing of the shampoo.

Without being bound by theory, it is believed that the coacervates provide
two major effects to the present shampoo composition. First, it reduces the
Critical Micelle Concentration (hereinafter "CMC") of the composition. The
reduction of the CMC relates to reduction of the surface tension, thereby
705 improving lather performance. Second, the existence of the anionic surfactants
along with the polyvalent metal cations expand the coacervate region in the
composition. As the cationic conditioning agents in the composition are mainly
delivered to the hair via these coacervates, expansion of the coacervate region
results in delivery of more cationic conditioning agents to the hair. Consequently,

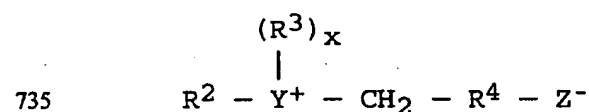
710 compositions which both cleanse and condition the hair from a single product, which have improved overall conditioning benefits and improved lathering are provided.

Techniques of analysis of formation of complex coacervates are known in the art. For example, microscopic analysis of the shampoo compositions, at any
715 chosen stage of dilution, can be utilized to identify whether a coacervate phase has formed. Such coacervate phase will be identifiable as an additional emulsified phase in the composition. The use of dyes can aid in distinguishing the coacervate phase from other insoluble phases dispersed in the shampoo composition.

720 Amphoteric And Zwitterionic Surfactant

Amphoteric surfactants for use herein include the derivatives of aliphatic secondary and tertiary amines in which the aliphatic radical is straight or branched and one of the aliphatic substituents contains from about 8 to about 18 carbon atoms and one contains an anionic water solubilizing group, e.g.,
725 carboxy, sulfonate, sulfate, phosphate, or phosphonate.

Zwitterionic surfactants for use herein include the derivatives of aliphatic quaternary ammonium, phosphonium, and sulfonium compounds, in which the aliphatic radicals are straight or branched, and wherein one of the aliphatic substituents contains from about 8 to about 18 carbon atoms and one contains
730 an anionic group, e.g., carboxy, sulfonate, sulfate, phosphate, or phosphonate. A general formula for these compounds is:

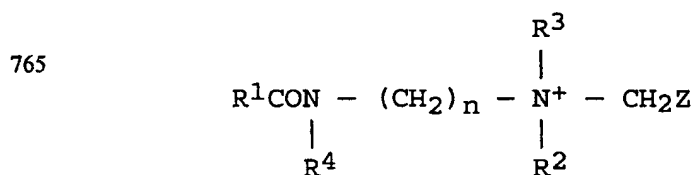


where R^2 contains an alkyl, alkenyl, or hydroxy alkyl radical of from about 8 to about 18 carbon atoms, from 0 to about 10 ethylene oxide moieties and from 0 to about 1 glyceryl moiety; Y is selected from the group consisting of nitrogen, phosphorus, and sulfur atoms; R^3 is an alkyl or monohydroxyalkyl group
740 containing 1 to about 3 carbon atoms; X is 1 when Y is a sulfur atom, and 2 when Y is a nitrogen or phosphorus atom; R^4 is an alkylene or hydroxyalkylene of from 1 to about 4 carbon atoms and Z is a radical selected from the group consisting of carboxylate, sulfonate, sulfate, phosphonate, and phosphate groups.

745 Examples of amphoteric and zwitterionic surfactants also include sultaines and amidosultaines. Sultaines, including amidosultaines, include for example, cocodimethylpropylsultaine, stearyldimethylpropylsultaine, lauryl-bis-(2-hydroxyethyl)propylsultaine and the like; and the amidosultaines such as cocamidodimethylpropylsultaine, stearylamidodimethylpropylsultaine, 750 laurylamido-bis-(2-hydroxyethyl)propylsultaine, and the like. Preferred are amidohydroxysultaines such as the C₈-C₁₈ hydrocarbylamidopropyl hydroxysultaines, especially C₈-C₁₄ hydrocarbylamidopropylhydroxysultaines, e.g., laurylamidopropylhydroxysultaine and cocamidopropylhydroxysultaine. Other sultaines are described in U.S. Patent 3,950,417, which is incorporated 755 herein by reference in its entirety.

Other suitable amphoteric surfactants are the aminoalkanoates of the formula RNH(CH₂)_nCOOM, the iminodialkanoates of the formula RN[(CH₂)_mCOOM]₂ and mixtures thereof; wherein n and m are numbers from 1 to about 4, R is C₈ - C₂₂ alkyl or alkenyl, and M is hydrogen, alkali metal, 760 alkaline earth metal, ammonium or alkanolammonium.

Other suitable amphoteric surfactants include those represented by the formula :



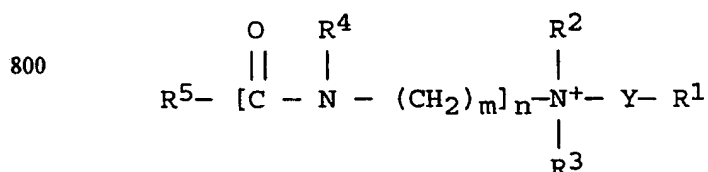
770 wherein R¹ is C₈ - C₂₂ alkyl or alkenyl, preferably C₈ - C₁₆. R² and R³ is independently selected from the group consisting of hydrogen, -CH₂CO₂M, -CH₂CH₂OH, -CH₂CH₂OCH₂CH₂COOM, or -(CH₂CH₂O)_mH wherein m is an integer from 1 to about 25, and R⁴ is hydrogen, -CH₂CH₂OH, or CH₂CH₂OCH₂CH₂COOM, Z is CO₂M or CH₂CO₂M, n is 2 or 3, preferably 2, M 775 is hydrogen or a cation, such as alkali metal (e.g., lithium, sodium, potassium), alkaline earth metal (beryllium, magnesium, calcium, strontium, barium), or ammonium. This type of surfactant is sometimes classified as an imidazoline-type amphoteric surfactant, although it should be recognized that it does not necessarily have to be derived, directly or indirectly, through an imidazoline 780 intermediate. Suitable materials of this type are marketed under the tradename MIRANOL and are understood to comprise a complex mixture of species, and

can exist in protonated and non-protonated species depending upon pH with respect to species that can have a hydrogen at R². All such variations and species are meant to be encompassed by the above formula.

785 Examples of surfactants of the above formula are monocarboxylates and di-carboxylates. Examples of these materials include cocoamphocarboxypropionate, cocoamphocarboxypropionic acid, cocoamphocarboxyglycinate (alternately referred to as cocoamphodiacetate), and cocoamphoacetate.

790 Commercial amphoteric surfactants include those sold under the trade names MIRANOL C2M CONC. N.P., MIRANOL C2M CONC. O.P., MIRANOL C2M SF, MIRANOL CM SPECIAL (Miranol, Inc.); ALKATERIC 2CIB (Alkaril Chemicals); AMPHOTERGE W-2 (Lonza, Inc.); MONATERIC CDX-38, MONATERIC CSH-32 (Mona Industries); REWOTERIC AM-2C (Rewo Chemical Group); and SCHERCOTERIC MS-2 (Scher Chemicals).

795 Betaine surfactants, i.e. zwitterionic surfactants, suitable for use in the conditioning compositions are those represented by the formula:



805 wherein: R¹ is a member selected from the group consisting of

COOM and CH(OH)CH₂SO₃M

R² is lower alkyl or hydroxyalkyl; R³ is lower alkyl or hydroxyalkyl; R⁴ is a member selected from the group consisting of hydrogen and lower alkyl; R⁵ is higher alkyl or alkenyl; Y is lower alkyl, preferably methyl; m is an integer from 2 to 7, preferably from 2 to 3; n is the integer 1 or 0; M is hydrogen or a cation, as previously described, such as an alkali metal, alkaline earth metal, or ammonium. The term "lower alkyl" or "hydroxyalkyl" means straight or branch chained, saturated, aliphatic hydrocarbon radicals and substituted hydrocarbon radicals having from one to about three carbon atoms such as, for example, methyl, ethyl, propyl, isopropyl, hydroxypropyl, hydroxyethyl, and the like. The term "higher alkyl or alkenyl" means straight or branch chained saturated (i.e., "higher alkyl") and unsaturated (i.e., "higher alkenyl") aliphatic hydrocarbon radicals having from about 8 to about 20 carbon atoms such as, for example, lauryl, cetyl, stearyl,

820 oleyl, and the like. It should be understood that the term "higher alkyl or alkenyl" includes mixtures of radicals which may contain one or more intermediate linkages such as ether or polyether linkages or non-functional substituents such as hydroxyl or halogen radicals wherein the radical remains of hydrophobic character.

825 Examples of surfactant betaines of the above formula wherein n is zero which are useful herein include the alkylbetaines such as cocodimethylcarboxymethylbetaine, lauryldimethylcarboxymethylbetaine, lauryldimethyl- α -carboxyethylbetaine, cetyldimethylcarboxymethylbetaine, lauryl-bis-(2-hydroxyethyl)-carboxymethylbetaine, stearyl-bis-(2-hydroxypropyl) carboxymethylbetaine, oleyldimethyl- γ -carboxypropylbetaine, lauryl-bis-(2-
830 hydroxypropyl)- α -carboxyethylbetaine, etc. The sulfobetaines may be represented by cocodimethylsulfopropylbetaine, stearyldimethylsulfopropyl betaine, lauryl-bis-(2-hydroxyethyl)-sulfopropylbetaine, and the like.

Specific examples of amido betaines and amidosulfobetaines useful in the conditioning compositions include the amidocarboxybetaines, such as cocamido
835 dimethylcarboxymethylbetaine, laurylamidodimethylcarboxymethylbetaine, cetylamidodimethylcarboxymethylbetaine, laurylamido-bis-(2-hydroxyethyl) carboxymethylbetaine, cocamido-bis-(2-hydroxyethyl)-carboxymethylbetaine, etc. The amidosulfobetaines may be represented by cocamidodimethyl sulfopropylbetaine, stearylamidodimethylsulfopropylbetaine, laurylamido-bis-(2-
840 hydroxyethyl)-sulfopropylbetaine, and the like.

Nonionic Surfactant

The compositions of the present invention can comprise a nonionic surfactant. Nonionic surfactants include those compounds produced by
845 condensation of alkylene oxide groups, hydrophilic in nature, with an organic hydrophobic compound, which may be aliphatic or alkyl aromatic in nature.

Preferred nonlimiting examples of nonionic surfactants for use in the shampoo compositions include the following:

(1) polyethylene oxide condensates of alkyl phenols, e.g., the
850 condensation products of alkyl phenols having an alkyl group containing from about 6 to about 20 carbon atoms in either a straight chain or branched chain configuration, with ethylene oxide, the said ethylene oxide being present in amounts equal to from about 10 to about 60 moles of ethylene oxide per mole of alkyl phenol;

(2) those derived from the condensation of ethylene oxide with the product
855 resulting from the reaction of propylene oxide and ethylene diamine products;

(3) condensation products of aliphatic alcohols having from about 8 to
about 18 carbon atoms, in either straight chain or branched chain configurations,
with ethylene oxide, e.g., a coconut alcohol ethylene oxide condensate having
from about 10 to about 30 moles of ethylene oxide per mole of coconut alcohol,
860 the coconut alcohol fraction having from about 8 to about 14 carbon atoms;

(4) long chain tertiary amine oxides of the formula $[R^1R^2R^3N \rightarrow O]$
where R^1 contains an alkyl, alkenyl or monohydroxy alkyl radical of from about 8
to about 18 carbon atoms, from 0 to about 10 ethylene oxide moieties, and from
0 to about 1 glyceryl moiety, and R^2 and R^3 contain from about 1 to about 3
865 carbon atoms and from 0 to about 1 hydroxy group, e.g., methyl, ethyl, propyl,
hydroxyethyl, or hydroxypropyl radicals;

(5) long chain tertiary phosphine oxides of the formula $[RR'R''P \rightarrow O]$
where R contains an alkyl, alkenyl or monohydroxyalkyl radical ranging from
about 8 to about 18 carbon atoms in chain length, from 0 to about 10 ethylene
oxide moieties and from 0 to 1 glyceryl moieties and R' and R'' are each alkyl or
870 monohydroxyalkyl groups containing from about 1 to about 3 carbon atoms;

(6) long chain dialkyl sulfoxides containing one short chain alkyl or
hydroxy alkyl radical of from 1 to about 3 carbon atoms (usually methyl) and one
long hydrophobic chain which include alkyl, alkenyl, hydroxy alkyl, or keto alkyl
875 radicals containing from about 8 to about 20 carbon atoms, from 0 to about 10
ethylene oxide moieties and from 0 to 1 glyceryl moieties;

(7) alkyl polysaccharide (APS) surfactants (e.g. alkyl polyglycosides),
examples of which are described in U.S. Patent 4,565,647, which is incorporated
herein by reference in its entirety, and which discloses APS surfactants having a
880 hydrophobic group with about 6 to about 30 carbon atoms and a polysaccharide
(e.g., polyglycoside) as the hydrophilic group; optionally, there can be a
polyalkylene-oxide group joining the hydrophobic and hydrophilic moieties; and
the alkyl group (i.e., the hydrophobic moiety) can be saturated or unsaturated,
branched or unbranched, and unsubstituted or substituted (e.g., with hydroxy or
885 cyclic rings); a preferred material is alkyl polyglucoside which is commercially
available from Henkel, ICI Americas, and Seppic; and

(8) polyoxyethylene alkyl ethers such as those of the formula
 $RO(CH_2CH_2)_nH$ and polyethylene glycol (PEG) glyceryl fatty esters, such as

those of the formula $R(O)OCH_2CH(OH)CH_2(OCH_2CH_2)_nOH$, wherein n is from
 890 1 to about 200, preferably from about 20 to about 100, and R is an alkyl having
 from about 8 to about 22 carbon atoms.

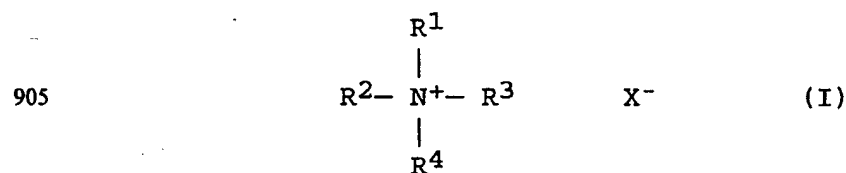
CONDITIONING AGENTS

Conditioning agents useful herein include cationic surfactants, high
 melting point compounds, oily compounds, cationic polymers, silicone
 895 compounds, and nonionic polymers. Conditioning agents may be present in the
 compositions herein either in combination with deterative surfactant, or without
 deterative surfactant.

Cationic Surfactant

The cationic surfactants useful herein are any known to the artisan.

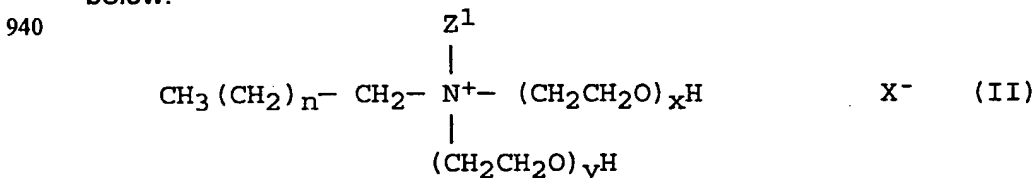
900 Among the cationic surfactants useful herein are those corresponding to
 the general formula (I):



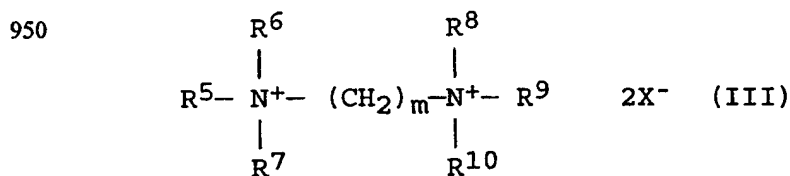
wherein at least one of R^1 , R^2 , R^3 , and R^4 is selected from an aliphatic group of
 910 from 8 to 30 carbon atoms or an aromatic, alkoxy, polyoxyalkylene, alkylamido,
 hydroxyalkyl, aryl or alkylaryl group having up to about 22 carbon atoms, the
 remainder of R^1 , R^2 , R^3 , and R^4 are independently selected from an aliphatic
 group of from 1 to about 22 carbon atoms or an aromatic, alkoxy,
 polyoxyalkylene, alkylamido, hydroxyalkyl, aryl or alkylaryl group having up to
 915 about 22 carbon atoms; and X is a salt-forming anion such as those selected
 from halogen, (e.g. chloride, bromide), acetate, citrate, lactate, glycolate,
 phosphate, nitrate, sulfonate, sulfate, alkylsulfate, and alkyl sulfonate radicals.
 The aliphatic groups can contain, in addition to carbon and hydrogen atoms,
 ether linkages, and other groups such as amino groups. The longer chain
 920 aliphatic groups, e.g., those of about 12 carbons, or higher, can be saturated or
 unsaturated. Preferred is when R^1 , R^2 , R^3 , and R^4 are independently selected
 from C_1 to about C_{22} alkyl. Nonlimiting examples of cationic surfactants useful
 include the materials having the following CTFA designations: quaternium-8,
 quaternium-24, quaternium-26, quaternium-27, quaternium-30, quaternium-33,
 925 quaternium-43, quaternium-52, quaternium-53, quaternium-56, quaternium-60,

quaternium-62, quaternium-70, quaternium-72, quaternium-75, quaternium-77, quaternium-78, quaternium-80, quaternium-81, quaternium-82, quaternium-83, quaternium-84, and mixtures thereof.

Also preferred are hydrophilically substituted cationic surfactants in which
 930 at least one of the substituents contain one or more aromatic, ether, ester, amido, or amino moieties present as substituents or as linkages in the radical chain, wherein at least one of the R¹ - R⁴ radicals contain one or more hydrophilic moieties selected from alkoxy (preferably C₁ - C₃ alkoxy), polyoxyalkylene (preferably C₁ - C₃ polyoxyalkylene), alkylamido, hydroxyalkyl,
 935 alkylester, and combinations thereof. Preferably, the hydrophilically substituted cationic conditioning surfactant contains from 2 to about 10 nonionic hydrophile moieties located within the above stated ranges. Preferred hydrophilically substituted cationic surfactants include those of the formula (II) through (VII) below:

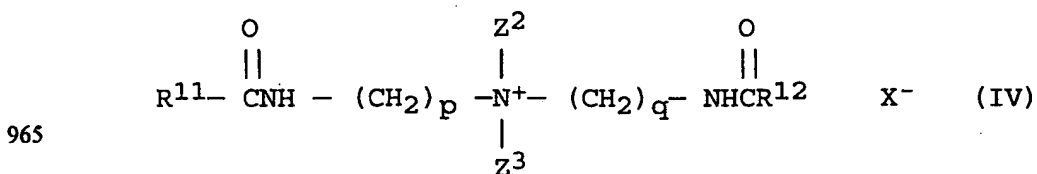


945 wherein n is from 8 to about 28, x+y is from 2 to about 40, Z¹ is a short chain alkyl, preferably a C₁ - C₃ alkyl, more preferably methyl, or - (CH₂CH₂O)_zH wherein x+y+z is up to 60, and X is a salt forming anion as defined above;

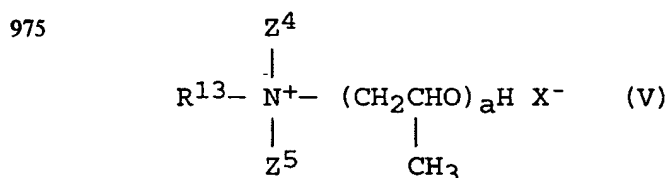


955 wherein m is 1 to 5, one or more of R⁵, R⁶, and R⁷ are independently an C₁ - C₃₀ alkyl, the remainder are - CH₂CH₂OH, one or two of R⁸, R⁹, and R¹⁰ are independently an C₁ - C₃₀ alkyl, and remainder are - CH₂CH₂OH, and X is a salt forming anion as mentioned above;

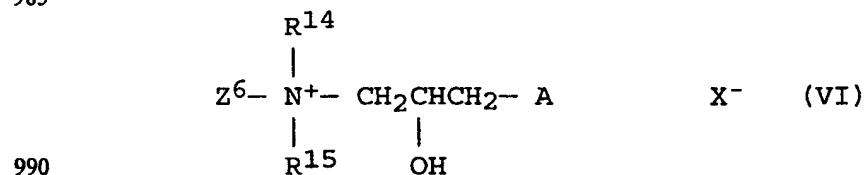
960



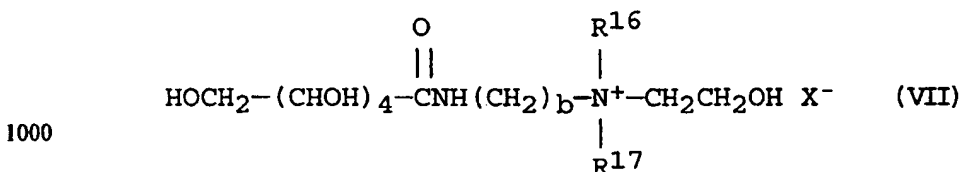
970 wherein Z^2 is an alkyl, preferably a C_1 - C_3 alkyl, more preferably methyl, and Z^3 is a short chain hydroxyalkyl, preferably hydroxymethyl or hydroxyethyl, p and q independently are integers from 2 to 4, inclusive, preferably from 2 to 3, inclusive, more preferably 2, R^{11} and R^{12} , independently, are substituted or unsubstituted hydrocarbyls, preferably C_{12} - C_{20} alkyl or alkenyl, and X is a salt forming anion as defined above;



980 wherein R^{13} is a hydrocarbyl, preferably a C_1 - C_3 alkyl, more preferably methyl, Z^4 and Z^5 are, independently, short chain hydrocarbyls, preferably C_2 - C_4 alkyl or alkenyl, more preferably ethyl, a is from 2 to about 40, preferably from about 7 to about 30, and X is a salt forming anion as defined above;



995 wherein R^{14} and R^{15} , independently, are C_1 - C_3 alkyl, preferably methyl, Z^6 is a C_{12} - C_{22} hydrocarbyl, alkyl carboxy or alkylamido, and A is a protein, preferably a collagen, keratin, milk protein, silk, soy protein, wheat protein, or hydrolyzed forms thereof; and X is a salt forming anion as defined above;



wherein b is 2 or 3, R¹⁶ and R¹⁷, independently are C₁ - C₃ hydrocarbyls preferably methyl, and X is a salt forming anion as defined above. Nonlimiting
1005 examples of hydrophilically substituted cationic surfactants useful include the materials having the following CTFA designations: quaternium-16, quaternium-61, quaternium-71, quaternium-79 hydrolyzed collagen, quaternium-79 hydrolyzed keratin, quaternium-79 hydrolyzed milk protein, quaternium-79 hydrolyzed silk, quaternium-79 hydrolyzed soy protein, and quaternium-79
1010 hydrolyzed wheat protein. Highly preferred compounds include commercially available materials of the following tradenames; VARIQUAT K1215 and 638 from Witco Chemical, MACKPRO KLP, MACKPRO WLW, MACKPRO MLP, MACKPRO NSP, MACKPRO NLW, MACKPRO WWP, MACKPRO NLP, MACKPRO SLP from McIntyre, ETHOQUAD 18/25, ETHOQUAD O/12PG, ETHOQUAD C/25, ETHOQUAD S/25, and ETHODUOQUAD from Akzo,
1015 DEHYQUAT SP from Henkel, and ATLAS G265 from ICI Americas.

Salts of primary, secondary, and tertiary fatty amines are also suitable cationic surfactants. The alkyl groups of such amines preferably have from about 12 to about 22 carbon atoms, and can be substituted or unsubstituted.
1020 Particularly useful are amido substituted tertiary fatty amines. Such amines, useful herein, include stearamidopropyldimethylamine, stearamidopropyldiethylamine, stearamidoethyldimethylamine, palmitamidopropyldimethylamine, palmitamidopropyldiethylamine, palmitamidoethyldimethylamine, behenamidopropyldimethylamine, behenamidopropyldiethylamine, behenamidoethyldimethylamine, arachidamidopropyldimethylamine, arachidamidopropyldiethylamine, arachidamidoethyldimethylamine, diethylaminoethylstearamide. Also useful are
1030 dimethylstearamine, dimethylsoyamine, soyamine, myristylamine, tridecylamine, ethylstearylamine, N-tallowpropane diamine, ethoxylated (with 5 moles of ethylene oxide) stearylamine, dihydroxyethylstearylamine, and arachidylbehenylamine. These amines can also be used in combination with acids such as L-glutamic acid, lactic acid, hydrochloric acid, malic acid, succinic
1035 acid, acetic acid, fumaric acid, tartaric acid, citric acid, L-glutamic hydrochloride, and mixtures thereof; more preferably L-glutamic acid, lactic acid, citric acid. Cationic amine surfactants included among those useful are disclosed in U.S.

Patent 4,275,055, Nachtigal, et al., issued June 23, 1981, which is incorporated by reference herein in its entirety.

1040 The cationic surfactants for use herein may also include a plurality of ammonium quaternary moieties or amino moieties, or a mixture thereof.

High Melting Point Compound

1045 The compositions may comprise a high melting point compound having a melting point of at least about 25°C selected from the group consisting of fatty alcohols, fatty acids, fatty alcohol derivatives, fatty acid derivatives, hydrocarbons, steroids, and mixtures thereof. Without being bound by theory, it is believed that these high melting point compounds cover the hair surface and reduce friction, thereby resulting in providing smooth feel on the hair and ease of combing. It is understood by the artisan that the compounds disclosed in this section of the specification can in some instances fall into more than one classification, e.g., some fatty alcohol derivatives can also be classified as fatty acid derivatives. However, a given classification is not intended to be a limitation on that particular compound, but is done so for convenience of classification and nomenclature. Further, it is understood by the artisan that, depending on the number and position of double bonds, and length and position of the branches, certain compounds having certain required carbon atoms may have a melting point of less than about 25°C. Such compounds of low melting point are not intended to be included in this section. Nonlimiting examples of the high melting point compounds are found in International Cosmetic Ingredient Dictionary, Fifth Edition, 1993, and CTFA Cosmetic Ingredient Handbook, Second Edition, 1992, both of which are incorporated by reference herein in their entirety.

1055 The fatty alcohols useful herein are those having from about 14 to about 30 carbon atoms, preferably from about 16 to about 22 carbon atoms. These fatty alcohols can be straight or branched chain alcohols and can be saturated or unsaturated. Nonlimiting examples of fatty alcohols include, cetyl alcohol, stearyl alcohol, behenyl alcohol, and mixtures thereof.

1065 The fatty acids useful herein are those having from about 10 to about 30 carbon atoms, preferably from about 12 to about 22 carbon atoms, and more preferably from about 16 to about 22 carbon atoms. These fatty acids can be straight or branched chain acids and can be saturated or unsaturated. Also included are diacids, triacids, and other multiple acids which meet the requirements herein. Also included herein are salts of these fatty acids.

1070

Nonlimiting examples of fatty acids include lauric acid, palmitic acid, stearic acid, behenic acid, sebacic acid, and mixtures thereof.

1075 The fatty alcohol derivatives and fatty acid derivatives useful herein include alkyl ethers of fatty alcohols, alkoxylated fatty alcohols, alkyl ethers of alkoxylated fatty alcohols, esters of fatty alcohols, fatty acid esters of compounds having esterifiable hydroxy groups, hydroxy-substituted fatty acids, and mixtures thereof. Nonlimiting examples of fatty alcohol derivatives and fatty acid
1080 derivatives include materials such as methyl stearyl ether; the ceteth series of compounds such as ceteth-1 through ceteth-45, which are ethylene glycol ethers of cetyl alcohol, wherein the numeric designation indicates the number of ethylene glycol moieties present; the steareth series of compounds such as steareth-1 through 10, which are ethylene glycol ethers of steareth alcohol, wherein the numeric designation indicates the number of ethylene glycol moieties present; cetareth 1 through cetareth-10, which are the ethylene glycol ethers of cetareth alcohol, i.e. a mixture of fatty alcohols containing predominantly cetyl and stearyl alcohol, wherein the numeric designation indicates the number of ethylene glycol moieties present; C₁-C₃₀ alkyl ethers of the ceteth, steareth, and
1085 cetareth compounds just described; polyoxyethylene ethers of behenyl alcohol; ethyl stearate, cetyl stearate, cetyl palmitate, stearyl stearate, myristyl myristate, polyoxyethylene cetyl ether stearate, polyoxyethylene stearyl ether stearate, polyoxyethylene lauryl ether stearate, ethyleneglycol monostearate, polyoxyethylene monostearate, polyoxyethylene distearate, propyleneglycol
1090 monostearate, propyleneglycol distearate, trimethylolpropane distearate, sorbitan stearate, polyglyceryl stearate, glyceryl monostearate, glyceryl distearate, glyceryl tristearate, and mixtures thereof.

Hydrocarbons useful herein include compounds having at least about 20 carbons.

1100 Steroids useful herein include compounds such as cholesterol.

High melting point compounds of a single compound of high purity are preferred. Single compounds of pure fatty alcohols selected from the group of pure cetyl alcohol, stearyl alcohol, and behenyl alcohol are highly preferred. By "pure" herein, what is meant is that the compound has a purity of at least about
1105 90%, preferably at least about 95%. These single compounds of high purity provide good rinsability from the hair when the consumer rinses off the composition.

Commercially available high melting point compounds useful herein include: cetyl alcohol, stearyl alcohol, and behenyl alcohol having tradenames
1110 KONOL series available from New Japan Chemical (Osaka, Japan), and NAA series available from NOF (Tokyo, Japan); pure behenyl alcohol having tradename 1-DOCOSANOL available from WAKO (Osaka, Japan), various fatty acids having tradenames NEO-FAT available from Akzo (Chicago Illinois, USA), HYSTRENE available from Witco Corp. (Dublin Ohio, USA), and DERMA
1115 available from Vevy (Genova, Italy); and cholesterol having tradename NIKKOL AGUASOME LA available from Nikko.

Oily Compound

The compositions comprise an oily compound having a melting point of not more than about 25°C selected from the group consisting of a first oily
1120 compound, a second oily compound, and mixtures thereof. The oily compounds useful herein may be volatile or nonvolatile. Without being bound by theory, it is believed that, the oily compounds may penetrate the hair to modify the hydroxy bonds of the hair, thereby resulting in providing softness and flexibility to the hair. The oily compound may comprise either the first oily compound or the second
1125 oily compound as described herein. Preferably, a mixture of the first oily compound and the second oily compound is used. The oily compounds of this section are to be distinguished from the high melting point compounds described above. Nonlimiting examples of the oily compounds are found in International Cosmetic Ingredient Dictionary, Fifth Edition, 1993, and CTFA Cosmetic
1130 Ingredient Handbook, Second Edition, 1992, both of which are incorporated by reference herein in their entirety.

First Oily Compound

The fatty alcohols useful herein include those having from about 10 to about 30 carbon atoms, preferably from about 12 to about 22 carbon atoms, and
1135 more preferably from about 16 to about 22 carbon atoms. These fatty alcohols can be straight or branched chain alcohols and can be saturated or unsaturated alcohols, preferably unsaturated alcohols. Nonlimiting examples of these compounds include oleyl alcohol, palmitoleic alcohol, isostearyl alcohol, isocetyl alcohol, undecanol, octyl dodecanol, octyl decanol, octyl alcohol, caprylic alcohol,
1140 decyl alcohol and lauryl alcohol.

The fatty acids useful herein include those having from about 10 to about 30 carbon atoms, preferably from about 12 to about 22 carbon atoms, and more

preferably from about 16 to about 22 carbon atoms. These fatty acids can be straight or branched chain acids and can be saturated or unsaturated. Suitable
1145 fatty acids include, for example, oleic acid, linoleic acid, isostearic acid, linolenic acid, ethyl linolenic acid, ethyl linolenic acid, arachidonic acid, and ricinolic acid.

The fatty acid derivatives and fatty alcohol derivatives are defined herein to include, for example, esters of fatty alcohols, alkoxylated fatty alcohols, alkyl
1150 ethers of fatty alcohols, alkyl ethers of alkoxylated fatty alcohols, and mixtures thereof. Nonlimiting examples of fatty acid derivatives and fatty alcohol derivatives, include, for example, methyl linoleate, ethyl linoleate, isopropyl linoleate, isodecyl oleate, isopropyl oleate, ethyl oleate, octyldodecyl oleate, oleyl oleate, decyl oleate, butyl oleate, methyl oleate, octyldodecyl stearate, octyldodecyl isostearate, octyldodecyl isopalmitate, octyl isopelargonate, octyl
1155 pelargonate, hexyl isostearate, isopropyl isostearate, isodecyl isononanoate, Oleth-2, pentaerythritol tetraoleate, pentaerythritol tetraisostearate, trimethylolpropane trioleate, and trimethylolpropane triisostearate.

Commercially available first oily compounds useful herein include: oleyl alcohol with tradename UNJECOL 90BHR available from New Japan Chemical,
1160 pentaerythritol tetraisostearate and trimethylolpropane triisostearate with tradenames KAKPTI and KAKTTI available from Kokyu Alcohol (Chiba, Japan), pentaerythritol tetraoleate having the same tradename as the compound name available from New Japan Chemical, trimethylolpropane trioleate with a tradename ENUJERUBU series available from New Japan Chemical, various
1165 liquid esters with tradenames SCHERCEMOL series available from Scher, and hexyl isostearate with a tradename HIS and isopropyl isostearate having a tradename ZPIS available from Kokyu Alcohol.

Second Oily Compound

The second oily compounds useful herein include straight chain, cyclic,
1170 and branched chain hydrocarbons which can be either saturated or unsaturated, so long as they have a melting point of not more than about 25°C. These hydrocarbons have from about 12 to about 40 carbon atoms, preferably from about 12 to about 30 carbon atoms, and preferably from about 12 to about 22 carbon atoms. Also encompassed herein are polymeric hydrocarbons of alkenyl
1175 monomers, such as polymers of C₂₋₆ alkenyl monomers. These polymers can be straight or branched chain polymers. The straight chain polymers will typically be relatively short in length, having a total number of carbon atoms as described

above. The branched chain polymers can have substantially higher chain lengths. The number average molecular weight of such materials can vary
1180 widely, but will typically be up to about 500, preferably from about 200 to about 400, and more preferably from about 300 to about 350. Also useful herein are the various grades of mineral oils. Mineral oils are liquid mixtures of hydrocarbons that are obtained from petroleum. Specific examples of suitable hydrocarbon materials include paraffin oil, mineral oil, dodecane, isododecane,
1185 hexadecane, isohexadecane, eicosene, isoeicosene, tridecane, tetradecane, polybutene, polyisobutene, and mixtures thereof. Preferred for use herein are hydrocarbons selected from the group consisting of mineral oil, isododecane, isohexadecane, polybutene, polyisobutene, and mixtures thereof.

Commercially available second oily compounds useful herein include
1190 isododecane, isohexadecane, and isoeicosene with tradenames PERMETHYL 99A, PERMETHYL 101A, and PERMETHYL 1082, available from Presperse (South Plainfield New Jersey, USA), a copolymer of isobutene and normal butene with tradenames INDOPOL H-100 available from Amoco Chemicals (Chicago Illinois, USA), mineral oil with tradename BENOL available from Witco,
1195 isoparaffin with tradename ISOPAR from Exxon Chemical Co. (Houston Texas, USA), α -olefin oligomer with tradename PURESYN 6 from Mobil Chemical Co., and trimethylolpropane tricaprilate/tricaprate with tradename MOBIL ESTER P43 from Mobil Chemical Co.

Cationic Polymers

1200 As used herein, the term "polymer" shall include materials whether made by polymerization of one type of monomer or made by two (i.e., copolymers) or more types of monomers.

Preferably, the cationic polymer is a water-soluble cationic polymer. By "water soluble" cationic polymer, what is meant is a polymer which is sufficiently
1205 soluble in water to form a substantially clear solution to the naked eye at a concentration of 0.1% in water (distilled or equivalent) at 25°C. The preferred polymer will be sufficiently soluble to form a substantially clear solution at 0.5% concentration, more preferably at 1.0% concentration.

The cationic polymers hereof will generally have a weight average
1210 molecular weight which is at least about 5,000, typically at least about 10,000, and is less than about 10 million. Preferably, the molecular weight is from about 100,000 to about 2 million. The cationic polymers will generally have cationic

nitrogen-containing moieties such as quaternary ammonium or cationic amino moieties, and mixtures thereof.

1215 The cationic charge density is preferably at least about 0.1 meq/gram, more preferably at least about 1.5 meq/gram, even more preferably at least about 1.1 meq/gram, still more preferably at least about 1.2 meq/gram. Cationic charge density of the cationic polymer can be determined according to the Kjeldahl Method. Those skilled in the art will recognize that the charge density of
1220 amino-containing polymers may vary depending upon pH and the isoelectric point of the amino groups. The charge density should be within the above limits at the pH of intended use.

Any anionic counterions can be utilized for the cationic polymers so long as the water solubility criteria is met. Suitable counterions include halides (e.g.,
1225 Cl, Br, I, or F, preferably Cl, Br, or I), sulfate, and methylsulfate. Others can also be used, as this list is not exclusive.

The cationic nitrogen-containing moiety will be present generally as a substituent, on a fraction of the total monomer units of the cationic hair conditioning polymers. Thus, the cationic polymer can comprise copolymers,
1230 terpolymers, etc. of quaternary ammonium or cationic amine-substituted monomer units and other non-cationic units referred to herein as spacer monomer units. Such polymers are known in the art, and a variety can be found in the CTFA Cosmetic Ingredient Dictionary, 3rd edition, edited by Estrin, Crosley, and Haynes, (The Cosmetic, Toiletry, and Fragrance Association, Inc.,
1235 Washington, D.C., 1982).

Suitable cationic polymers include, for example, copolymers of vinyl monomers having cationic amine or quaternary ammonium functionalities with water soluble spacer monomers such as acrylamide, methacrylamide, alkyl and dialkyl acrylamides, alkyl and dialkyl methacrylamides, alkyl acrylate, alkyl
1240 methacrylate, vinyl caprolactone, and vinyl pyrrolidone. The alkyl and dialkyl substituted monomers preferably have C₁ - C₇ alkyl groups, more preferably C₁ - C₃ alkyl groups. Other suitable spacer monomers include vinyl esters, vinyl alcohol (made by hydrolysis of polyvinyl acetate), maleic anhydride, propylene glycol, and ethylene glycol.

1245 The cationic amines can be primary, secondary, or tertiary amines, depending upon the particular species and the pH of the composition. In general, secondary and tertiary amines, especially tertiary amines, are preferred.

Amine-substituted vinyl monomers can be polymerized in the amine form, and then optionally can be converted to ammonium by a quaternization reaction. 1250 Amines can also be similarly quaternized subsequent to formation of the polymer. For example, tertiary amine functionalities can be quaternized by reaction with a salt of the formula R'X wherein R' is a short chain alkyl, preferably a C₁ - C₇ alkyl, more preferably a C₁ - C₃ alkyl, and X is an anion which forms a water soluble salt with the quaternized ammonium.

1255 Suitable cationic amino and quaternary ammonium monomers include, for example, vinyl compounds substituted with dialkylaminoalkyl acrylate, dialkylaminoalkyl methacrylate, monoalkylaminoalkyl acrylate, monoalkylaminoalkyl methacrylate, trialkyl methacryloxyalkyl ammonium salt, trialkyl acryloxyalkyl ammonium salt, diallyl quaternary ammonium salts, and vinyl 1260 quaternary ammonium monomers having cyclic cationic nitrogen-containing rings such as pyridinium, imidazolium, and quaternized pyrrolidone, e.g., alkyl vinyl imidazolium, alkyl vinyl pyridinium, alkyl vinyl pyrrolidone salts. The alkyl portions of these monomers are preferably lower alkyls such as the C₁ - C₃ alkyls, more preferably C₁ and C₂ alkyls. Suitable amine-substituted vinyl monomers for use 1265 herein include dialkylaminoalkyl acrylate, dialkylaminoalkyl methacrylate, dialkylaminoalkyl acrylamide, and dialkylaminoalkyl methacrylamide, wherein the alkyl groups are preferably C₁ - C₇ hydrocarbyls, more preferably C₁ - C₃, alkyls.

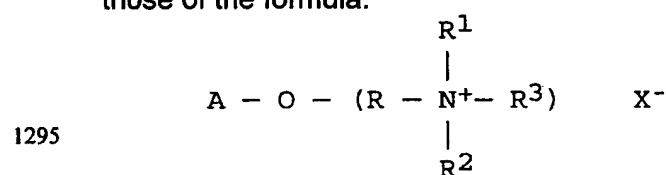
The cationic polymers hereof can comprise mixtures of monomer units 1270 derived from amine- and/or quaternary ammonium-substituted monomer and/or compatible spacer monomers.

Suitable cationic hair conditioning polymers include, for example: copolymers of 1-vinyl-2-pyrrolidone and 1-vinyl-3-methylimidazolium salt (e.g., chloride salt) (referred to in the industry by the Cosmetic, Toiletry, and Fragrance 1275 Association, "CTFA", as Polyquaternium-16), such as those commercially available from BASF Wyandotte Corp. (Parsippany, NJ, USA) under the LUVIQUAT tradename (e.g., LUVIQUAT FC 370); copolymers of 1-vinyl-2-pyrrolidone and dimethylaminoethyl methacrylate (referred to in the industry by CTFA as Polyquaternium-11) such as those commercially available from Gaf Corporation (Wayne, NJ, USA) under the GAFQUAT tradename (e.g., GAFQUAT 1280 755N); cationic diallyl quaternary ammonium-containing polymers, including, for example, dimethyldiallylammonium chloride homopolymer and copolymers of

acrylamide and dimethyldiallylammonium chloride, referred to in the industry (CTFA) as Polyquaternium 6 and Polyquaternium 7, respectively; and mineral
 1285 acid salts of amino-alkyl esters of homo- and co-polymers of unsaturated carboxylic acids having from 3 to 5 carbon atoms, as described in U.S. Patent 4,009,256, incorporated herein by reference.

Other cationic polymers that can be used include polysaccharide polymers, such as cationic cellulose derivatives and cationic starch derivatives.

1290 Cationic polysaccharide polymer materials suitable for use herein include those of the formula:



wherein: A is an anhydroglucose residual group, such as a starch or cellulose anhydroglucose residual, R is an alkylene oxyalkylene, polyoxyalkylene, or
 1300 hydroxyalkylene group, or combination thereof, R¹, R², and R³ independently are alkyl, aryl, alkylaryl, arylalkyl, alkoxyalkyl, or alkoxyaryl groups, each group containing up to about 18 carbon atoms, and the total number of carbon atoms for each cationic moiety (i.e., the sum of carbon atoms in R¹, R² and R³) preferably being about 20 or less, and X is an anionic counterion, as previously
 1305 described.

Cationic cellulose is available from Amerchol Corp. (Edison, NJ, USA) in their Polymer JR® and LR® series of polymers, as salts of hydroxyethyl cellulose reacted with trimethyl ammonium substituted epoxide, referred to in the industry (CTFA) as Polyquaternium 10. Another type of cationic cellulose includes the
 1310 polymeric quaternary ammonium salts of hydroxyethyl cellulose reacted with lauryl dimethyl ammonium-substituted epoxide, referred to in the industry (CTFA) as Polyquaternium 24. These materials are available from Amerchol Corp. (Edison, NJ, USA) under the tradename Polymer LM-200®.

Other cationic polymers that can be used include cationic guar gum
 1315 derivatives, such as guar hydroxypropyltrimonium chloride (commercially available from Celanese Corp. in their Jaguar R series). Other materials include quaternary nitrogen-containing cellulose ethers (e.g., as described in U.S. Patent 3,962,418, incorporated herein by reference), and copolymers of etherified

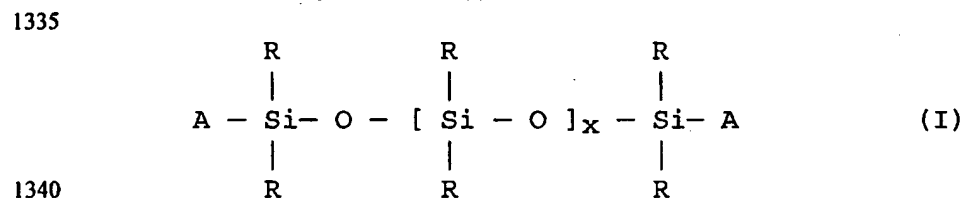
cellulose and starch (e.g., as described in U.S. Patent 3,958,581, incorporated
1320 herein by reference.)

Silicone Compounds

The conditioning agents useful herein include silicone compounds. The
silicone compounds hereof can include volatile soluble or insoluble, or nonvolatile
soluble or insoluble silicone conditioning agents. By soluble what is meant is that
1325 the silicone compound is miscible with the carrier of the composition so as to
form part of the same phase. By insoluble what is meant is that the silicone
forms a separate, discontinuous phase from the carrier, such as in the form of an
emulsion or a suspension of droplets of the silicone.

Suitable silicone fluids include polyalkyl siloxanes, polyaryl siloxanes,
1330 polyalkylaryl siloxanes, polyether siloxane copolymers, and mixtures thereof.
Other nonvolatile silicone compounds having hair conditioning properties can
also be used.

The silicone compounds herein also include polyalkyl or polyaryl siloxanes
with the following structure (I)



wherein R is alkyl or aryl, and x is an integer from about 7 to about 8,000. "A"
represents groups which block the ends of the silicone chains. The alkyl or aryl
groups substituted on the siloxane chain (R) or at the ends of the siloxane chains
1345 (A) can have any structure as long as the resulting silicone remains fluid at room
temperature, is dispersible, is neither irritating, toxic nor otherwise harmful when
applied to the hair, is compatible with the other components of the composition,
is chemically stable under normal use and storage conditions, and is capable of
being deposited on and conditions the hair. Suitable A groups include hydroxy,
1350 methyl, methoxy, ethoxy, propoxy, and aryloxy. The two R groups on the silicon
atom may represent the same group or different groups. Preferably, the two R
groups represent the same group. Suitable R groups include methyl, ethyl,
propyl, phenyl, methylphenyl and phenylmethyl. The preferred silicone
compounds are polydimethylsiloxane, polydiethylsiloxane, and
1355 polymethylphenylsiloxane. Polydimethylsiloxane, which is also known as

dimethicone, is especially preferred. The polyalkylsiloxanes that can be used include, for example, polydimethylsiloxanes. These silicone compounds are available, for example, from the General Electric Company in their ViscasilR and SF 96 series, and from Dow Corning in their Dow Corning 200 series.

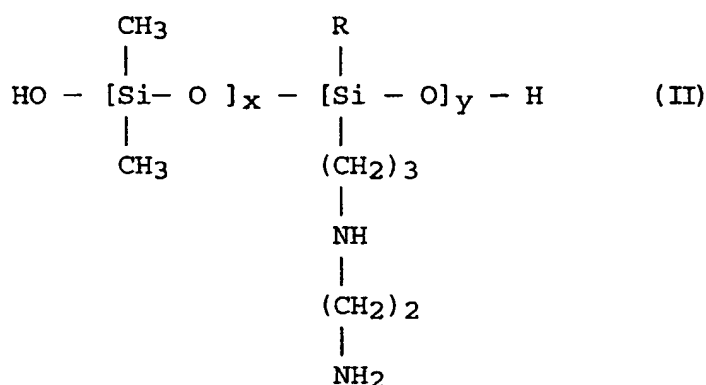
1360 Polyalkylaryl siloxane fluids can also be used and include, for example, polymethylphenylsiloxanes. These siloxanes are available, for example, from the General Electric Company as SF 1075 methyl phenyl fluid or from Dow Corning as 556 Cosmetic Grade Fluid.

1365 Especially preferred, for enhancing the shine characteristics of hair, are highly arylated silicone compounds, such as highly phenylated polyethyl silicone having refractive index of about 1.46 or higher, especially about 1.52 or higher. When these high refractive index silicone compounds are used, they should be mixed with a spreading agent, such as a surfactant or a silicone resin, as described below to decrease the surface tension and enhance the film forming
1370 ability of the material.

The silicone compounds that can be used include, for example, a polypropylene oxide modified polydimethylsiloxane although ethylene oxide or mixtures of ethylene oxide and propylene oxide can also be used. The ethylene oxide and polypropylene oxide level should be sufficiently low so as not to
1375 interfere with the dispersibility characteristics of the silicone. These material are also known as dimethicone copolyols.

Other silicone compounds include amino substituted materials. Suitable alkylamino substituted silicone compounds include those represented by the following structure (II)

1380



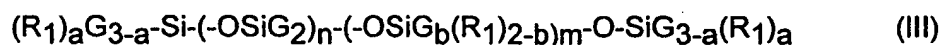
1385

1390

wherein R is CH₃ or OH, x and y are integers which depend on the molecular weight, the average molecular weight being approximately between 5,000 and 10,000. This polymer is also known as "amodimethicone".

1395

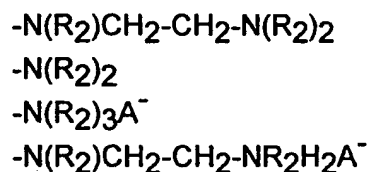
Suitable amino substituted silicone fluids include those represented by the formula (III)



1400

in which G is chosen from the group consisting of hydrogen, phenyl, OH, C₁-C₈ alkyl and preferably methyl; a denotes 0 or an integer from 1 to 3, and preferably equals 0; b denotes 0 or 1 and preferably equals 1; the sum n+m is a number from 1 to 2,000 and preferably from 50 to 150, n being able to denote a number from 0 to 1,999 and preferably from 49 to 149 and m being able to denote an integer from 1 to 2,000 and preferably from 1 to 10; R₁ is a monovalent radical of formula C_qH_{2q}L in which q is an integer from 2 to 8 and L is chosen from the groups

1405

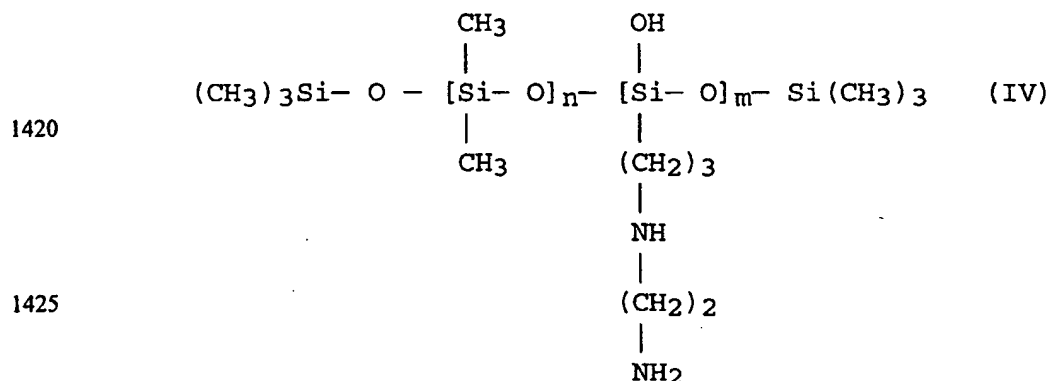


1410

in which R₂ is chosen from the group consisting of hydrogen, phenyl, benzyl, a saturated hydrocarbon radical, preferably an alkyl radical containing from 1 to 20 carbon atoms, and A⁻ denotes a halide ion.

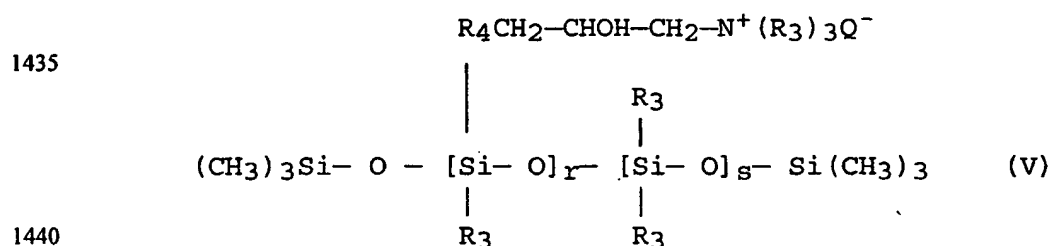
An especially preferred amino substituted silicone corresponding to formula (III) is the polymer known as "trimethylsilylamodimethicone", of formula (IV):

1415



1430 In this formula n and m are selected depending on the exact molecular weight of the compound desired.

Other amino substituted silicone polymers which can be used are represented by the formula (V):



1445 where R^3 denotes a monovalent hydrocarbon radical having from 1 to 18 carbon atoms, preferably an alkyl or alkenyl radical such as methyl; R_4 denotes a hydrocarbon radical, preferably a $\text{C}_1 - \text{C}_{18}$ alkylene radical or a $\text{C}_1 - \text{C}_{18}$, and more preferably $\text{C}_1 - \text{C}_8$, alkyleneoxy radical; Q^- is a halide ion, preferably chloride; r denotes an average statistical value from 2 to 20, preferably from 2 to 8; s denotes an average statistical value from 20 to 200, and preferably from 20 to 50. A preferred polymer of this class is available from Union Carbide under the name "UCAR SILICONE ALE 56."

1450 References disclosing suitable nonvolatile dispersed silicone compounds include U.S. Patent No. 2,826,551, to Geen; U.S. Patent No. 3,964,500, to Drakoff, issued June 22, 1976; U.S. Patent No. 4,364,837, to Pader; and British Patent No. 849,433, to Woolston, all of which are incorporated herein by reference in their entirety. Also incorporated herein by reference in its entirety is

1455 "Silicon Compounds" distributed by Petrarch Systems, Inc., 1984. This reference

provides an extensive, though not exclusive, listing of suitable silicone compounds.

Another nonvolatile dispersed silicone that can be especially useful is a silicone gum. The term "silicone gum", as used herein, means a polyorganosiloxane material having a viscosity at 25°C of greater than or equal to 1,000,000 centistokes. It is recognized that the silicone gums described herein can also have some overlap with the above-disclosed silicone compounds. This overlap is not intended as a limitation on any of these materials. Silicone gums are described by Petrarch, and others including U.S. Patent No. 4,152,416, to Spitzer et al., issued May 1, 1979 and Noll, Walter, Chemistry and Technology of Silicones, New York: Academic Press 1968. Also describing silicone gums are General Electric Silicone Rubber Product Data Sheets SE 30, SE 33, SE 54 and SE 76. All of these described references are incorporated herein by reference in their entirety. The "silicone gums" will typically have a mass molecular weight in excess of about 200,000, generally between about 200,000 and about 1,000,000. Specific examples include polydimethylsiloxane, poly(dimethylsiloxane methylvinylsiloxane) copolymer, poly(dimethylsiloxane diphenylsiloxane methylvinylsiloxane) copolymer and mixtures thereof.

Also useful are silicone resins, which are highly crosslinked polymeric siloxane systems. The crosslinking is introduced through the incorporation of tri-functional and tetra-functional silanes with mono-functional or di-functional, or both, silanes during manufacture of the silicone resin. As is well understood in the art, the degree of crosslinking that is required in order to result in a silicone resin will vary according to the specific silane units incorporated into the silicone resin. In general, silicone materials which have a sufficient level of trifunctional and tetrafunctional siloxane monomer units, and hence, a sufficient level of crosslinking, such that they dry down to a rigid, or hard, film are considered to be silicone resins. The ratio of oxygen atoms to silicon atoms is indicative of the level of crosslinking in a particular silicone material. Silicone materials which have at least about 1.1 oxygen atoms per silicon atom will generally be silicone resins herein. Preferably, the ratio of oxygen:silicon atoms is at least about 1.2:1.0. Silanes used in the manufacture of silicone resins include monomethyl-, dimethyl-, trimethyl-, monophenyl-, diphenyl-, methylphenyl-, monovinyl-, and methylvinylchlorosilanes, and tetrachlorosilane, with the methyl substituted silanes being most commonly utilized. Preferred resins are offered by General

Electric as GE SS4230 and SS4267. Commercially available silicone resins will generally be supplied in a dissolved form in a low viscosity volatile or nonvolatile silicone fluid. The silicone resins for use herein should be supplied and incorporated into the present compositions in such dissolved form, as will be readily apparent to those skilled in the art. Without being bound by theory, it is believed that the silicone resins can enhance deposition of other silicone compounds on the hair and can enhance the glossiness of hair with high refractive index volumes.

Other useful silicone resins are silicone resin powders such as the material given the CTFA designation polymethylsilsequioxane, which is commercially available as TospearlTM from Toshiba Silicones.

The method of manufacturing these silicone compounds, can be found in Encyclopedia of Polymer Science and Engineering, Volume 15, Second Edition, pp 204-308, John Wiley & Sons, Inc., 1989, which is incorporated herein by reference in its entirety.

Silicone materials and silicone resins in particular, can conveniently be identified according to a shorthand nomenclature system well known to those skilled in the art as the "MDTQ" nomenclature. Under this system, the silicone is described according to the presence of various siloxane monomer units which make up the silicone. Briefly, the symbol M denotes the mono-functional unit $(\text{CH}_3)_3\text{SiO}_{1.5}$; D denotes the difunctional unit $(\text{CH}_3)_2\text{SiO}$; T denotes the trifunctional unit $(\text{CH}_3)\text{SiO}_{1.5}$; and Q denotes the quadri- or tetra-functional unit SiO_2 . Primes of the unit symbols, e.g., M', D', T', and Q' denote substituents other than methyl, and must be specifically defined for each occurrence. Typical alternate substituents include groups such as vinyl, phenyl, amino, hydroxyl, etc. The molar ratios of the various units, either in terms of subscripts to the symbols indicating the total number of each type of unit in the silicone, or an average thereof, or as specifically indicated ratios in combination with molecular weight, complete the description of the silicone material under the MDTQ system. Higher relative molar amounts of T, Q, T' and/or Q' to D, D', M and/or M' in a silicone resin is indicative of higher levels of crosslinking. As discussed before, however, the overall level of crosslinking can also be indicated by the oxygen to silicon ratio.

The silicone resins for use herein which are preferred are MQ, MT, MTQ, MQ and MDTQ resins. Thus, the preferred silicone substituent is methyl.

Especially preferred are MQ resins wherein the M:Q ratio is from about 0.5:1.0 to about 1.5:1.0 and the average molecular weight of the resin is from about 1000 to about 10,000.

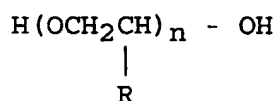
Nonionic Polymer

1530 Nonionic polymers useful herein include cellulose derivatives, hydrophobically modified cellulose derivatives, ethylene oxide polymers, and ethylene oxide/propylene oxide based polymers. Suitable nonionic polymers are cellulose derivatives including methylcellulose with tradename BENECEL, hydroxyethyl cellulose with tradename NATROSOL, hydroxypropyl cellulose with
1535 tradename KLUCEL, cetyl hydroxyethyl cellulose with tradename POLYSURF 67, all supplied by Hercules. Other suitable nonionic polymers are ethylene oxide and/or propylene oxide based polymers with tradenames CARBOWAX PEGs, POLYOX WASRs, and UCON FLUIDS, all supplied by Amerchol.

Polyalkylene Glycols

1540 These compounds are particularly useful for compositions which are designed to impart a soft, moist feeling to the hair. When present, the polyalkylene glycol is typically used at a level from about 0.025% to about 1.5%, preferably from about 0.05% to about 1%, and more preferably from about 0.1% to about 0.5% of the compositions.

1545 The polyalkylene glycols are characterized by the general formula:



1550 wherein R is selected from the group consisting of H, methyl, and mixtures thereof. When R is H, these materials are polymers of ethylene oxide, which are also known as polyethylene oxides, polyoxyethylenes, and polyethylene glycols. When R is methyl, these materials are polymers of propylene oxide, which are also known as polypropylene oxides, polyoxypropylenes, and polypropylene
1555 glycols. When R is methyl, it is also understood that various positional isomers of the resulting polymers can exist.

In the above structure, n has an average value of from about 1500 to about 25,000, preferably from about 2500 to about 20,000, and more preferably from about 3500 to about 15,000.

1560 Polyethylene glycol polymers useful herein are PEG-2M wherein R equals H and n has an average value of about 2,000 (PEG-2M is also known as Polyox WSR®).

N-10, which is available from Union Carbide and as PEG-2,000); PEG-5M wherein R equals H and n has an average value of about 5,000 (PEG-5M is also known as Polyox WSR® N-35 and Polyox WSR® N-80, both available from Union Carbide and as PEG-5,000 and Polyethylene Glycol 300,000); PEG-7M wherein R equals H and n has an average value of about 7,000 (PEG-7M is also known as Polyox WSR® N-750 available from Union Carbide); PEG-9M wherein R equals H and n has an average value of about 9,000 (PEG 9-M is also known as Polyox WSR® N-3333 available from Union Carbide); and PEG-14 M wherein R equals H and n has an average value of about 14,000 (PEG-14M is also known as Polyox WSR® N-3000 available from Union Carbide).

Other useful polymers include the polypropylene glycols and mixed polyethylene/polypropylene glycols.

ADDITIONAL COMPONENTS

The compositions of the present invention may include a variety of additional components, which may be selected by the artisan according to the desired characteristics of the final product. Additional components include, for example, polyvalent metal cations, suspending agents, and other additional components.

Polyvalent Metal Cations

Suitable polyvalent metal cations include divalent and trivalent metals, divalent metals being preferred. Exemplary metal cations include alkaline earth metals, such as magnesium, calcium, zinc, and copper, and trivalent metals such as aluminum and iron. Preferred are calcium and magnesium.

The polyvalent metal cation can be added as an inorganic salt, organic salt, or as a hydroxide. The polyvalent metal cation may also be added as a salt with anionic surfactants as mentioned above.

Preferably, the polyvalent metal cation is introduced as an inorganic salt or organic salt. Inorganic salts include chloride, bromide, iodine, nitrate, or sulfate, more preferably chloride or sulfate. Organic salts include L-glutamate, lactate, malate, succinate, acetate, fumarate, L-glutamic acid hydrochloride, and tartarate.

It will be clear to those skilled in the art that, if polyvalent salts of the anionic surfactant is used as the mode of introducing the polyvalent metal cations into the compositions hereof, only a fraction of the anionic surfactant may be of

polyvalent form, the remainder of the anionic surfactant being necessarily added in monovalent form.

Hardness of the conditioning shampoo compositions can be measured by standard methods in the art, such as by ethylene diamine tetraacetic acid (EDTA) titration. In the event that the composition contains dyes or other color materials that interfere with the ability of EDTA titration to yield a perceptible color change, hardness should be determined from the composition in the absence of the interfering dye or color.

Suspending Agents

A preferred additional component is a suspending agent, particularly for compositions comprising silicone compounds of high viscosity and/or large particle size. When present, the suspending agent is in dispersed form in the compositions. The suspending agent will generally comprise from about 0.1% to about 10%, and more typically from about 0.3% to about 5.0%, by weight, of the composition.

Preferred suspending agents include acyl derivatives such as ethylene glycol stearates, both mono and distearate, long chain amine oxides such as alkyl (C₁₆-C₂₂) dimethyl amine oxides, e.g., stearyl dimethyl amine oxide, and mixtures thereof. When used in the shampoo compositions, these preferred suspending agents are present in the composition in crystalline form. These suspending agents are described in U.S. Patent 4,741,855.

Other suitable suspending agents include alkanol amides of fatty acids, preferably having from about 16 to about 22 carbon atoms, more preferably about 16 to 18 carbon atoms, preferred examples of which include stearic monoethanolamide, cocomonethanolamide, stearic diethanolamide, stearic monoisopropanolamide and stearic monoethanolamide stearate.

Other suitable suspending agents include N,N-dihydrocarbyl amido benzoic acid and soluble salts thereof (e.g., Na and K salts), particularly N,N-di(hydrogenated) C₁₆, C₁₈ and tallow amido benzoic acid species of this family, which are commercially available from Stepan Company (Northfield, Illinois, USA).

Other suitable suspending agents include xanthan gum. The use of xanthan gum as a suspending agent in silicone containing shampoo compositions is described, for example, in U.S. Patent 4,788,006, which is incorporated herein by reference in its entirety. Combinations of long chain acyl

derivatives and xanthan gum may also be used as a suspending agent in the shampoo compositions. Such combinations are described in U.S. Patent 4,704,272, which is incorporated herein by reference in its entirety.

Other suitable suspending agents include carboxyvinyl polymers. Preferred among these polymers are the copolymers of acrylic acid crosslinked with polyallylsucrose as described in U.S. Patent 2,798,053, which is incorporated herein by reference in its entirety. Examples of these polymers include the carbomers, which are homopolymers of acrylic acid crosslinked with an allyl ether of pentaerythritol, an allyl ether of sucrose, or an allyl ether of propylene. Neutralizers may be required, for example, amino methyl propanol, triethanol amine, or sodium hydroxide.

Other suitable suspending agents can be used in the compositions, including those that can impart a gel-like viscosity to the composition, such as water soluble or colloidal water soluble polymers like cellulose ethers such as hydroxyethyl cellulose, hydroxymethyl cellulose, hydroxypropyl cellulose, and materials such as guar gum, polyvinyl alcohol, polyvinyl pyrrolidone, hydroxypropyl guar gum, starch and starch derivatives.

Other Additional Components

A wide variety of other additional ingredients can be formulated into the present compositions. These include: other conditioning agents such as hydrolysed collagen with tradename Peptin 2000 available from Hormel, vitamin E with tradename Emix-d available from Eisai, panthenol available from Roche, panthenyl ethyl ether available from Roche, hydrolysed keratin, proteins, plant extracts, and nutrients; hair-fixative polymers such as amphoteric fixative polymers, cationic fixative polymers, anionic fixative polymers, nonionic fixative polymers, and silicone grafted copolymers; preservatives such as benzyl alcohol, methyl paraben, propyl paraben and imidazolidinyl urea; pH adjusting agents, such as citric acid, sodium citrate, succinic acid, phosphoric acid, sodium hydroxide, sodium carbonate; salts, in general, such as potassium acetate and sodium chloride; coloring agents, such as any of the FD&C or D&C dyes; hair oxidizing (bleaching) agents, such as hydrogen peroxide, perborate and persulfate salts; hair reducing agents such as the thioglycolates; perfumes; and sequestering agents, such as disodium ethylenediamine tetra-acetate; ultraviolet and infrared screening and absorbing agents such as octyl salicylate, and antidandruff agents such as zinc pyridinethione. Such optional ingredients generally

are used individually at levels from about 0.001% to about 10.0%, preferably from about 0.01% to about 5.0% by weight of the composition.

EXAMPLES

1670 The following examples further describe and demonstrate embodiments within the scope of the present invention. The examples are given solely for the purpose of illustration and are not to be construed as limitations of the present invention, as many variations thereof are possible without departing from the spirit and scope of the invention. Ingredients are identified by chemical or CTFA name, or otherwise defined below.

Method of Preparation

The hair spray, mousses, lotions and conditioners are suitably made as follows: If included in the formula, polymeric materials are dispersed in water at room temperature. If required, the polymeric materials such as Carbomer and
1680 Acrylates/Steareth-20 Methacrylate Copolymer may be neutralized after dispersing. The mixture is then heated up to above 60°C, and fatty alcohols and emulsifiers are added if included in the formulation. After cooling down to below 50°C, the remaining components are added with agitation then cooled down to about 30°C. Ethanol is added here if included in the formula. A triblender and mill
1685 can be used if necessary to disperse the materials. As appropriate, the mixture thus obtained can be packed into an aerosol can with propellant.

The shampoos are suitably prepared by any conventional method well known in the art. A suitable method is as follows: polymer and surfactants are dispersed in water to form a homogeneous mixture. To this mixture are added the
1690 other ingredients except for silicone emulsion (if present), perfume, and salt; the obtained mixture is agitated. If present, the silicone emulsion is made with Dimethicone or Dimethiconol, a small amount of deterative surfactant, and a portion of water. The obtained mixture is then passed through a heat exchanger to cool, and the silicone emulsion, perfume, and salt are added. The obtained
1695 compositions are poured into bottles to make hair shampoo compositions.

Alternatively, water and surfactants and any other solids that need to be melted can be mixed together at elevated temperature, e.g., above about 70°C, to speed the mixing into shampoo. Additional ingredients can be added either to this hot premix or after cooling the premix. The ingredients are mixed thoroughly
1700 at the elevated temperature and then pumped through a high shear mill and then through a heat exchanger to cool them to ambient temperature. If present in the

composition, silicone emulsified at room temperature in concentrated surfactant is added to the cooled mix.

1705 Compositions

	Ex. 1	Ex.2	Ex.3	Ex.4	Ex.5	Ex.6
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	1.0	-	-	-	-	-
4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)-amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids *2	-	0.8	-	-	-	-
4-methyl-6,7-dihydroxycoumarin *3	-	-	1.0	-	-	-
4-methyl-7,7'-dimethylamino coumarin *4	-	-	-	0.1	-	-
2-(4-styryl-3-sulfophenyl)-2H-naphtho[1,2-d]triazole	-	-	-	1.0	-	-
1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline *5	-	-	-	-	0.5	-
2,4-dimethoxy-6-(1'pyrenyl)-1,3,5-triazine *6	-	-	-	-	-	1.0
Glycerin *7	-	-	5.0	-	-	-
Propylene Glycol *8	1.0	3.0	-	10.0	5.0	10.0
Ammonium Laureth-3 Sulfate	12.0	12.0	12.0	15.0	10.0	10.0
Ammonium Lauryl Sulfate	2.0	2.0	2.0	2.0	-	-
N-acyl-L-gultamate Triethanolamine *9	2.0	2.0	-	-	2.0	2.0
Sodium Lauroyl Sarcosinate *10	-	-	2.0	2.0	2.0	1.0
Cocamidopropylbetaine *11	2.0	4.0	4.0	4.0	2.0	2.0
Laureth-20 *12	-	0.5	-	0.5	0.5	-
Alkyl polyglucoside *13	1.0	-	1.0	-	-	-
Ethylene Glycol Distearate *14	2.0	2.0	2.0	2.0	2.0	1.5
Cocamide MEA	1.5	1.5	1.5	1.5	1.5	1.5
DMDM Hydantoin	0.2	0.2	0.2	0.2	0.2	0.2
Magnesium chloride	0.5	-	0.5	-	-	-
Silicone	3.0	2.0	2.0	3.0	1.6	2.0
Polyoxyethyleneglycol *16	-	0.2	-	0.1	0.5	-
Hydroxyethyl Cellulose *17	0.1	-	0.1	-	-	0.5

Polyquaternium-10 *18	0.5	0.4	-	-	-	-
Monoalkyl trimethyl ammonium	-	-	0.1	-	-	-
Cetyl alcohol *19	0.2	0.1	-	0.1	0.1	-
Stearyl alcohol *20	-	-	-	0.1	0.2	-
Behenyl alcohol *21	-	-	0.1	-	-	-
Perfume	0.5	0.5	0.5	0.5	0.5	0.5
Pentaerythritol Tetraisostearate *22	-	-	1.0	0.5	-	-
Trimethylolpropane Triisostearate *23	0.5	-	-	-	0.3	-
Pentaerythritol Tetraoleate	-	0.3	-	-	-	-
Trimethylolpropane Trioleate *24	-	-	-	-	-	0.5
Monosodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Disodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Hydrolyzed Collagen *39	0.01	-	-	-	-	-
Vitamin E *40	0.01	-	-	-	-	-
Panthenol *41	0.025	-	-	-	-	-
Panthenyl Ethyl Ether *42	0.225	-	-	-	-	-
Deionized Water	----- q.s. to 100% -----					

	Ex.7	Ex.8	Ex.9	Ex.10
Disodium-1,4-bis(2-sulfostyryl) biphenyl *1	0.2	1.0	0.5	0.8
Glycerin *7	3.0	-	5.0	-
Propylene Glycol *8	-	3.0	-	5.0
Ammonium Laureth-3 Sulfate	12.0	12.0	15.0	10.0
Ammonium Lauryl Sulfate	2.0	2.0	2.0	2.0
Sodium Lauroyl Sarcosinate *10	2.0	-	-	1.0
Cocamidopropylbetaine *11	1.0	2.0	-	2.0
Cocamide MEA	1.5	1.5	1.5	1.5
DMDM Hydantoin	0.2	0.2	0.2	0.2
Monosodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5
Disodium phosphate	0 to 0.5	0 to 0.5	0 to 0.5	0 to 0.5

Deionized Water	----- q.s. to 100% -----
-----------------	--------------------------

1710

	Ex.11	Ex.12	Ex.13	Ex.14	Ex.15
Disodium-1,4-bis(2-sulfoethyl)biphenyl *1	0.80	-	-	0.10	-
4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)amino-1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids *2	-	-	-	-	0.80
1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline *5	-	-	-	0.60	-
4,4'-bis(5-methylbenzoxazol-2-yl)stilbene *25	-	-	0.50	-	-
2,4-dimethoxy-6-(1'pyrenyl)-1,3,5-triazine *6	-	0.20	-	-	-
Glycerin *7	-	-	5.00	-	-
Propylene Glycol *8	-	-	-	5.00	-
Denatured Ethyl Alcohol	-	-	-	-	18.98
Cetyl Alcohol *19	2.00	0.50	0.60	0.20	0.60
Stearyl Alcohol *20	3.00	0.50	0.40	0.20	0.40
Behenyl Alcohol *21	-	-	-	0.20	-
Silicone Emulsion *15	1.00	0.50	2.00	0.20	1.00
Polysorbate 60 *26	1.00	-	-	-	-
Lauryl Methyl Gluceth-10 Hydroxypropyl-dimonium Chloride *27	-	1.00	-	0.50	-
Dihydrogenated Tallowamidoethyl Hydroxyethylmonium Methosulfate *28	-	-	0.10	-	0.10
Citric Acid *29	0-0.2	0-0.2	0-0.2	0-0.2	-
Hydroxyl Propyl Cellulose *30	-	-	-	-	0.50
Polyoxyethylene Glycol *16	0.50	0.50	0.50	0.50	0.50
Aminomethylpropanol *31	2.10	-	2.10	-	-
Acrylates/Steareth-20 Methacrylate Copolymer *32	1.75	-	1.75	-	-
Cetyl Hydroxyethyl Cellulose *33	-	0.75	-	0.75	-

Polyquaternium-10 *18	0.10	-	0.10	0.20	-
Polyquaternium-7 *34	-	0.50	-	-	1.00
Pentaerythritol Tetraisostearate *22	1.00	3.00	-	-	-
Trimethylolpropane Trioleate *24	-	-	0.50	-	0.10
Trimethylolpropane Triisostearate *23	-	-	-	0.20	-
Preservatives	0.90	0.90	0.90	0.90	-
Perfume	0.08	0.08	0.08	0.08	0.08
Deionized Water	----- q.s. to 100% -----				

	Ex.16	Ex.17	Ex.18	Ex.19
4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)amino 1,3,5-triazin-2-yl)amino]stilbene-2,2'-disulfonic acids *2	0.80	-	-	-
4-methyl-6,7-dihydroxycoumarin *3	-	0.50	-	-
4-methyl-7,7'-dimethylamino coumarin *4	-	-	0.50	-
2-(4-styryl-3-sulfophenyl)-2H-naphto[1,2-d]triazole	-	-	-	0.40
Glycerin *7	-	5.00	-	2.0
Propylene Glycol *8	-	-	5.00	-
Denatured Ethyl Alcohol	19.07	18.42	17.81	-
Cetyl Alcohol *19	0.20	-	2.00	2.50
Stearyl Alcohol *20	0.20	-	-	4.50
Behenyl Alcohol *21	0.10	0.50	-	-
15/85 Silicone Blend *35	-	-	-	4.20
Silicone Emulsion *15	0.20	0.50	1.00	
Stearamidopropyl dimethylamine *36	0.20	-	-	2.00
L-Glutamic Acid *34	0.08	-	-	0.64
Hydroxyethyl Cellulose *17	-	0.50	-	-
Polyoxyethylene Glycol *16	0.50	0.50	0.50	0.20

Carbomer *38	0.50	-	0.50	-
Aminomethylpropanol *31	0.60	-	0.60	-
Polyquaternium-10 *18	1.00	0.10	-	0.20
Polyquaternium-7 *34	-	-	0.50	-
Pentaerythritol Tetraisostearate *22	0.10	0.10	0.20	-
Trimethylolpropane Trioleate *24	0.10	0.10	0.05	-
Trimethylolpropane Triisostearate *23	-	-	-	0.25
Preservatives	-	-	-	0.53
Perfume	0.08	0.08	0.08	0.20
Deionized Water	----- q.s. to 100% -----			

	Ex. 20	Ex.21	Ex.22	Ex.23	Ex.24	Ex.25
Disodium-1,4-bis(2-sulfostyryl)biphenyl *1	0.20	0.20	0.20	1.60	0.80	1.60
Glycerin *7	3.00	-	15.00	-	3.00	-
Propylene Glycol *8	-	3.00	-	15.00	-	5.00
Denatured Ethyl Alcohol	-	-	-	-	19.22	18.66
Preservatives	0.90	0.90	0.90	0.90	-	-
Perfume	0.08	0.08	0.08	0.08	0.08	0.08
Deionized Water	----- q.s. to 100% -----					

Definitions

- 1715 *1 Disodium-1,4-bis(2-sulfostyryl)biphenyl: TINOPAL CBX obtained by Ciba Geigy.
- *2 4,4'-bis[(4-anilino-6-bis(2-hydroxyethyl)amino-1,3,5-triazin-2-yl)amino] stilbene-2,2'-disulfonic acids: TINOPAL UNPA-GX obtained by Ciba Geigy.
- *3 4-methyl-6,7-dihydroxycoumarin: available from Wako.
- *4 4-methyl-7,7'-dimethylamino coumarin: available from Wako.
- 1720 *5 1-(4-amidosulfonylphenyl)-3-(4-chlorophenyl)-2-pyrazoline: BLANKOPHOR DCB obtained by Bayer.
- *6 2,4-dimethoxy-6-(1'pyrenyl)-1,3,5-triazine: available from Ciba Geigy.
- *7 Glycerin: GLYCERIN USP obtained by Nihon Yushi.
- *8 Propylene Glycol: LEXOL PG-865 obtained by Inolex.

- *9 N-acyl-L-glutamate Triethanolamine: CT12S obtained by Ajinomoto.
- *10 Sodium Lauroyl Sarcosinate: SOYPON obtained by Kawaken Fine Chem.
- *11 Cocamidopropylbetaine: TEGOBETAIN obtained by Th.Goldschmidt AG.
- 1730 *12 Laureth-20: BL-20 obtained by Nikko.
- *13 Alkyl polyglucoside: PLANTACARE 2000UP obtained by Henkel.
- *14 Ethylene Glycol Distearate: EGDS obtained by Th. Goldschmidt AG.
- *15 Silicone Emulsion: X65-4829 obtained by Tosil/GE.
- *16 Polyoxyethylene Glycol: WSR N-10 obtained by Amerchol.
- 1735 *17 Hydroxyethyl Cellulose: available from Aqualon.
- *18 Polyquaternium-10: UCARE POLYMER LR 400 obtained by Amerchol.
- *19 Cetyl Alcohol: KONOL SERIES obtained by Shinihon Rika.
- *20 Stearyl Alcohol: KONOL SERIES obtained by Shinihon Rika.
- *21 Behenyl Alcohol: 1-DOCOSANOL (97%) obtained by Wako.
- 1740 *22 Pentaerythritol Tetraisostearate: KAK PTI obtained by Kokyu alcohol.
- *23 Trimethylolpropane Triisostearate: KAK TTI obtained by Kokyu alcohol.
- *24 Trimethylolpropane Trioleate: ENJERUBU TP3SO obtained by Shinihon Rika.
- *25 4,4'-bis(5-methylbenzoxazol-2-yl)stilbene: available from TCI.
- 1745 *26 Polysorbate 60: TWEEN 60 obtained by ICI.
- *27 Lauryl Methyl Gluceth-10 Hydroxypropyldimonium Chloride: GLUCQUAT 125 obtained by Amerchol.
- *28 Dihydrogenated Tallowamidoethyl Hydroxyethylmonium Methosulfate: VARISOFT 110
- 1750 obtained by Witco.
- *29 Citric Acid: ANHYDROUS CITRIC ACID obtained by Haarman & Reimer.
- *30 Hydroxyl Propyl Cellulose: METLOSE obtained by Shinetsu.
- *31 Aminomethylpropanol: AMP-REGULAR obtained by Angus.
- *32 Acrylates/Steareth-20 Methacrylate Copolymer: ACULYN obtained by 125
- 1755 Rohm&Haas.
- *33 Cetyl Hydroxyethyl Cellulose: POLYSURF 67 obtained by Aqualon.
- *34 Polyquaternium-7: MERQUAT S obtained by Calgon.
- *35 15/85 Silicone Blend: available from Shinetsu.
- *36 Stearamidopropyl dimethylamine: AMIDOAMINE MPS obtained by Nikko.
- 1760 *37 L-Glutamic Acid: L-GLUTAMIC ACID (COSMETIC GRADE) obtained by Ajinomoto.

- *38 Carbomer: CARBOPOL 980 obtained by BF Goodrich.
- *39 Hydrolyzed Collagen: PEPTEIN 2000 obtained by Hormel.
- *40 Vitamin E: EMIX-d obtained by Eisai.
- 1765 *41 Panthenol: available from Roche.
- *42 Panthenyl Ethyl Ether: available from Roche.

1770 The embodiments disclosed and represented by the previous examples have many advantages. For example, they can provide the benefits associated with optical brighteners (e.g., shine, UV protection, color alteration) with enhanced deposition of the optical brightener on the hair, prevention of dulling or dusting of the hair due to crystallization of the optical brightener without reduction in the level of optical brightener used in the composition.

1775 It is understood that the examples and embodiments described herein are for illustrative purposes only and that various modifications or changes in light thereof will be suggested to one skilled in the art without departing from its spirit and scope.

WHAT IS CLAIMED IS:

1. A hair care composition comprising:
 - (a) an effective amount of an optical brightener; and
 - (b) a non-volatile solvent for solubilizing at least 0.01 wt% of the optical brightener;5 wherein the optical brightener is deposited on the hair without crystallizing.
2. The hair care composition of claim 1 further comprising a carrier suitable for application to the hair.
3. The hair care composition of claim 2 further comprising a detergent surfactant suitable for cleansing the hair.
4. The hair care composition of claim 3 further comprising a conditioning agent.
5. The hair care composition of claim 1 further comprising a conditioning agent.
6. The hair care composition of any of the preceding claims wherein the optical brightener is selected from the group consisting of polystyrylstilbenes, triazinstilbenes, hydroxycoumarins, aminocoumarins, triazoles, pyrazolines, oxazoles, pyrenes, porphyrins, imidazoles, and mixtures thereof.
5
7. The hair care composition of claim 6 wherein the non-volatile solvent is selected from the group consisting of alkyl alcohols, propane diol, propylene glycol, hexylene glycol, glycerin, diethylene glycol, dipropylene glycol, butylene glycol, and mixtures thereof.
5
8. The hair care composition of claim 7 containing at least about 0.1 wt% of the non-volatile solvent.

9. The hair care composition of claim 8 containing from about 0.001 wt% to about 20 wt% of the optical brightener.

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/16578

A. CLASSIFICATION OF SUBJECT MATTER IPC 6 A61K7/13 A61K7/00		
According to International Patent Classification (IPC) or to both national classification and IPC		
B. FIELDS SEARCHED Minimum documentation searched (classification system followed by classification symbols) IPC 6 A61K		
Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched		
Electronic data base consulted during the international search (name of data base and, where practical, search terms used)		
C. DOCUMENTS CONSIDERED TO BE RELEVANT		
Category *	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 746 864 A (GOLGATE PALMOLIVE-PEET CO) 21 March 1956 see page 2, line 59 - line 65 see page 3, line 5 - line 24 see page 3, line 49 see page 3, line 85 - line 88 see claims 1-12 ---	1,3,6-9
X	US 3 658 985 A (OLSON FRANK WESLEY JR ET AL) 25 April 1972 see column 1, line 63 - line 72 see column 2, line 1 - line 52 see column 3, line 55 - line 66 see column 4, line 67 - line 70 see column 5; example 2 see claims --- -/---	1-12
<div style="display: flex; justify-content: space-between;"> <input checked="" type="checkbox"/> Further documents are listed in the continuation of box C. <input checked="" type="checkbox"/> Patent family members are listed in annex. </div>		
<div style="display: flex;"> <div style="flex: 1;"> <p>* Special categories of cited documents:</p> <p>"A" document defining the general state of the art which is not considered to be of particular relevance</p> <p>"E" earlier document but published on or after the international filing date</p> <p>"L" document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)</p> <p>"O" document referring to an oral disclosure, use, exhibition or other means</p> <p>"P" document published prior to the international filing date but later than the priority date claimed</p> </div> <div style="flex: 1;"> <p>"T" later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention</p> <p>"X" document of particular relevance; the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone</p> <p>"Y" document of particular relevance; the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art.</p> <p>"&" document member of the same patent family</p> </div> </div>		
Date of the actual completion of the international search <div style="text-align: center; font-weight: bold;">29 June 1998</div>		Date of mailing of the international search report <div style="text-align: center; font-weight: bold;">06/07/1998</div>
Name and mailing address of the ISA European Patent Office, P.B. 5818 Patentlaan 2 NL - 2280 HV Rijswijk Tel. (+31-70) 340-2040, Tx. 31 651 epo nl, Fax: (+31-70) 340-3016		Authorized officer <div style="text-align: center; font-weight: bold;">Pelli Wablat, B</div>

INTERNATIONAL SEARCH REPORT

International Application No
PCT/US 97/16578

C.(Continuation) DOCUMENTS CONSIDERED TO BE RELEVANT		
Category	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
X	GB 2 307 639 A (GEN ELECTRIC) 4 June 1997 see page 3, line 4 - line 10 see page 4, line 13 - line 17 see page 6, line 17 - line 34 see page 7, line 6 - line 35 see claims ---	1,2,6-9
X	US 4 126 674 A (MAUSNER JACK J) 21 November 1978 see column 2, line 28 - line 44 see example 2 ---	1-9
X	US 4 938 954 A (GROSS PAUL ET AL) 3 July 1990 see abstract see example 5 ---	1,2,6,7, 9
X	US 3 577 528 A (EDMAN WALTER W ET AL) 4 May 1971 see example 7 ---	1,4-6
X	EP 0 087 060 A (FABER CASTELL A W) 31 August 1983 cited in the application see example 5 -----	1,6,7

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/16578

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
GB 746864 A		BE 522974 A FR 1094336 A	18-05-1955
US 3658985 A	25-04-1972	AT 302534 A BE 753955 A CH 531884 A DE 2034295 A DK 125620 B FR 2053194 A GB 1307644 A NL 7011176 A SE 351564 B ZA 7004430 A	15-09-1972 31-12-1970 31-12-1972 11-02-1971 19-03-1973 16-04-1971 21-02-1973 01-02-1971 04-12-1972 23-02-1972
GB 2307639 A	04-06-1997	DE 19646804 A FR 2741261 A JP 9183714 A	22-05-1997 23-05-1997 15-07-1997
US 4126674 A	21-11-1978	AR 218450 A AU 508026 B AU 2504877 A BR 7702988 A CA 1091585 A DE 2721278 A FR 2350835 A JP 1391716 C JP 53004010 A JP 61054767 B ZA 7702805 A	13-06-1980 06-03-1980 16-11-1978 20-12-1977 16-12-1980 24-11-1977 09-12-1977 23-07-1987 14-01-1978 25-11-1986 26-04-1978
US 4938954 A	03-07-1990	DE 3725080 A DE 3864936 A WO 8900845 A EP 0301197 A JP 2515149 B JP 2500105 T	09-02-1989 24-10-1991 09-02-1989 01-02-1989 10-07-1996 18-01-1990
US 3577528 A	04-05-1971	DE 1617808 A GB 1152972 A NL 6703673 A	08-04-1971 21-05-1969 28-11-1967

INTERNATIONAL SEARCH REPORT

Information on patent family members

International Application No

PCT/US 97/16578

Patent document cited in search report	Publication date	Patent family member(s)	Publication date
EP 0087060 A	31-08-1983	DE 3204636 A JP 58198410 A	25-08-1983 18-11-1983
<hr/>			